



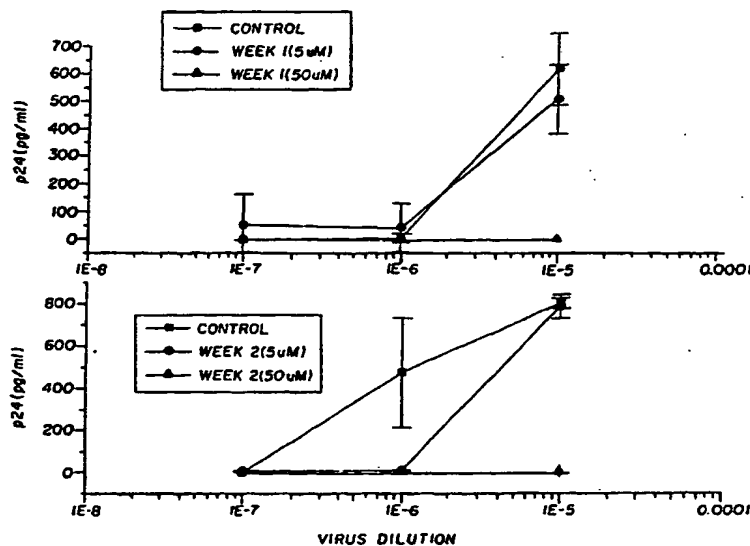
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(54) Title: STABLE COPPER(I) COMPLEXES AND METHODS RELATED THERETO



(57) Abstract

There is disclosed stable copper(I) complexes and methods relating thereto. The stable copper(I) complexes comprise a copper(I) ion complexed by a multi-dentate ligand which favors the +1 oxidation state for copper. Methods of this invention include the use of the stable copper(I) complexes as wound healing agents, anti-oxidative agents, anti-inflammatory agents, lipid modulating agents, signal transduction modulating agents, hair growth agents, and anti-viral agents. Exemplary stable copper(I) complexes include neocuproine copper(I) and bathocuproine disulfonic acid copper(I).

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DescriptionSTABLE COPPER(I) COMPLEXES  
AND METHODS RELATED THERETO

5

Technical Field

This invention is generally directed to a copper(I) complex and methods relating to the use thereof and, more specifically, to copper(I) complexed by a multi-dentate  
10 ligand such that the +1 oxidation state for copper is favored in the resulting complex.

Background of the Invention

Copper is found in both plants and animals, and a  
15 number of copper-containing proteins, including enzymes, have been isolated. Copper may exist in a variety of oxidation states, including the 0, +1, +2 and +3 oxidation states (i.e., copper(0), copper(I), copper(II) and copper(III), respectively), with copper(I) and copper(II)  
20 the most common. The relative stabilities of copper(I) and copper(II) in aqueous solution depend on the nature of the anions or other ligands present in the solution. Moreover, only low equilibrium concentrations of copper(I) in aqueous solutions (i.e.,  $< 10^{-2}M$ ) can exist. This  
25 instability is due, in part, to the tendency of copper(I) to disproportionate to copper(II) and copper(0). Most copper(I) compounds readily oxidize to copper(II) compounds, although further oxidation to copper(III) is difficult (see, generally, A.F. Cotton and G. Wilkinson,  
30 Advanced Inorganic Chemistry, 5th ed., John Wiley & Sons, New York, pp. 903-922, 1988).

Due to the relatively well-defined aqueous chemistry of copper(II), a large number of copper(II) salts and complexes are known. For example, a great deal of  
35 research has been directed to the biological activity of peptide/copper(II) complexes, and such copper(II)

complexes have been shown to possess utility for a variety of therapeutic and cosmetic purposes. In particular, the naturally occurring glycyl-histidyl-lysine:copper(II) complex ("GHK-Cu(II)") has been shown to be an effective agent in the enhancement of wound healing in warm-blooded animals, as well as generally serving as an anti-inflammatory agent (see U.S. Patent No. 4,760,051). Various derivatives of GHK-Cu(II) possess similar activity (see U.S. Patent Nos. 4,665,054 and 4,877,770). GHK-Cu(II) and other peptide-copper(II) complexes have also been shown to be effective for stimulating hair growth (U.S. Patent Nos. 5,177,061 and 5,120,831), for inducing biological coverings in wounds (U.S. Patent No. 4,810,693), for preventing ulcers (U.S. Patent Nos. 4,767,753, 5,023,237, 5,145,838), for cosmetic applications (U.S. Patent No. 5,135,913), and for healing bone (U.S. Patent No. 5,509,588). Moreover, anti-oxidative and anti-inflammatory activity of metal(II)-peptide complexes has been disclosed (U.S. Patent No. 5,118,665), as well as the use of copper(II)-containing compounds to accelerate wound healing (U.S. Patent No. 5,164,367).

Although great strides have been made in the study of copper(II) complexes, and particularly peptide/copper(II) complexes, there is still a need in the art for additional copper complexes which possess biological activity. The present invention fulfills this need, and provides further related advantages.

### Summary of the Invention

This invention is generally directed to stable copper(I) complexes and methods relating thereto. More specifically, the stable copper(I) complexes of the present invention comprise copper(I) complexed by a multidentate ligand such that the +1 oxidation state for copper is favored.

The stable copper(I) complexes have utility for enhancing wound healing in warm-blooded animals, for enhancing or restoring the resistance of warm-blooded animals to oxidative or inflammatory damage associated with reactive oxygen species and/or lipid mediators, for stimulating the growth of hair in warm-blooded animals, for modulating lipid metabolism, for modulating signal transduction in cells by inhibiting protein kinases, and for inhibiting viral activity, including (but not limited to) HIV replication in an HIV-infected animal. Methods of the present invention comprise administering an effective amount of a stable copper(I) complex to the animal.

Other aspects of this invention will become evident upon reference to the attached figures and following detailed description. All references identified in the detailed description, including the examples, are hereby incorporated by reference in their entirety

#### Description of the Figures

Figure 1 illustrates the activity of a representative copper(I) complex of this invention (i.e., bathocuproine disulfonic acid ("BCDS") copper(I)) to accelerate wound healing.

Figure 2 illustrates the ability of a representative copper(I) complex of the present invention, BCDS copper(I), to inhibit viral (i.e., HIV) replication.

Figure 3 illustrates synthesis pathways for prostaglandins and leukotrienes, as well as certain key enzymes associated therewith.

Figure 4 illustrates a synthesis pathway for cholesterol formation, including the intermediates acetyl CoA and HMG-CoA and the enzymes acetyl CoA synthetase and HMG-CoA reductase.

Figure 5 illustrates the action of Protein Kinase C (PKC) and protein tyrosine kinase in signal transduction (PI = phosphatidyl inositol, IP<sup>3</sup> = inositol triphosphate,

PG = phosphatyl glycerol, P-Protein = phosphorylated protein, CDR PK = calmodulin-regulated protein kinase, PKA = Protein Kinase A, Protein Kinase = Protein Tyrosine Kinase (cytoplasmic), and EGF-R Protein Kinase = Epidermal growth factor receptor protein tyrosine kinase).

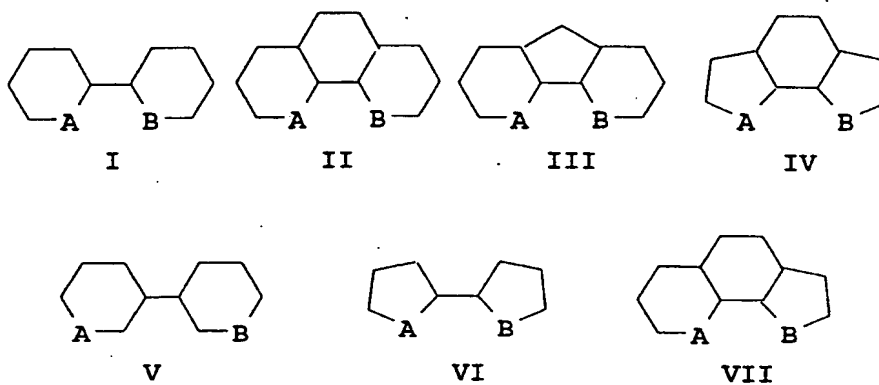
#### Detailed Description

This invention is generally directed to copper(I) complexes and methods relating to the use thereof, and more specifically, to copper(I) complexed by a multi-dentate ligand to form a stable copper(I) complex. As used herein, a "stable copper(I) complex" is copper(I) chelated by at least one multi-dentate ligand such that the resulting complex favors the +1 oxidation state of copper. The most common states of copper(I) are associated with four coordination sites, and are generally of a tetrahedral configuration. In general, chelating agents are coordination compounds in which a single ligand occupies more than one coordination position of a metal ion. If the ligand occupies two coordination positions, it is considered a bi-dentate ligand; if more than two coordination positions are occupied by the ligand, it is considered a poly-dentate ligand (such as a tri-dentate ligand or a tetra-dentate ligand). As used herein, a "multi-dentate ligand" is a bi-, tri- or tetra-dentate ligand which occupies two, three or four coordination sites, respectively, of copper (I).

The stable copper(I) complexes of this invention include all complexes of copper(I) chelated by at least one multi-dentate ligand which structurally favors the +1 oxidation state of copper. Copper(I) complexes may be formed by reacting a multi-dentate ligand with a source of copper(I) (such as CuCl, Cu<sub>2</sub>O or CuCN) in aqueous solution. The resulting copper(I) complex may then be observed by suitable analytical techniques, such as ESR, NMR and/or UV-VIS, to determine the oxidation state of the copper in

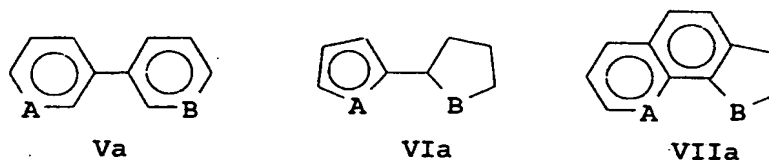
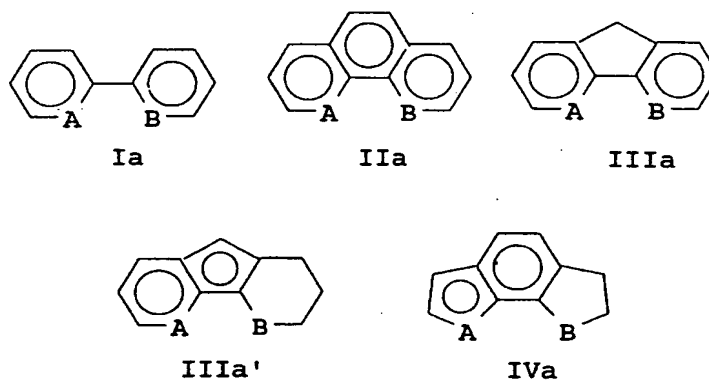
the complex (see Munakata et al., Copper Coordination Chemistry: Biochemical and Inorganic Perspectives, Karlin and Zubieta editors, Adenine Press, Guilderland, N.Y., pp. 473-495, 1983). For example, copper(I) complexes can be identified by their characteristic absence of an ESR signal, while copper(II) complexes will generally possess an ESR signal. Furthermore, copper(II) complexes exhibit broadening of proton NMR signals, and copper(I) complexes exhibit relatively sharp proton NMR signals. Following identification of the copper(I) complex, its stability can be evaluated by determining its susceptibility to oxidation by, for example, exposing the copper(I) complex to air. As used herein, a "stable" copper(I) complex has a half-life of at least 5 minutes, preferably of at least one hour, and more preferably of 24 hours or more (i.e., half of the copper(I) complex remains in the +1 oxidation state) upon exposure to air, at room temperature (23°C) and atmospheric pressure. In other words, stable copper(I) complexes of this invention resist oxidation, while non-stable copper(I) complexes are readily oxidized to yield copper(II) complexes upon exposure to air.

As mentioned above, any multi-dentate ligand which chelates copper(I) to yield a stable copper(I) complex is suitable in the practice of this invention. However, in a preferred embodiment, the multi-dentate ligands of this invention are selected from the following general structures I through VII:



wherein A and B represent heteroatoms which may occupy  
coordination sites of copper(I), and are preferably  
selected from nitrogen, oxygen, sulfur and phosphorous.

The rings of structures I through VII may be  
aromatic, non-aromatic or a mixture of both aromatic and  
non-aromatic rings. For example, the following structures  
are representative of such combinations:

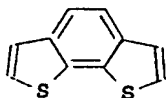
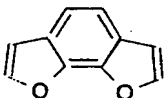
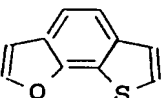
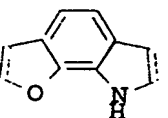
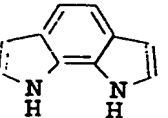


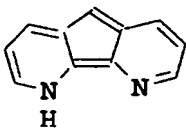
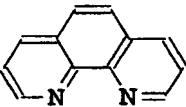
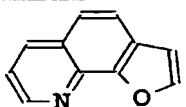
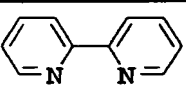


Representative examples of multi-dentate ligands of this invention having structures I through VII are set forth in Table 1. Specifically, Table 1 identifies the structure of the representative multi-dentate ligand, lists the corresponding chemical name, identifies the Chemical Abstracts Registration Number ("CA Reg. No."), and provides a corresponding reference (if available) describing the synthesis and/or chemistry of the identified multi-dentate ligand.

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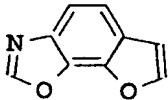
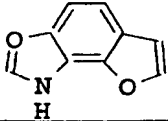
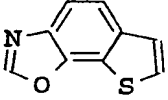
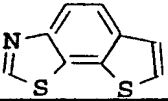
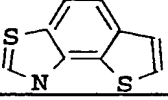
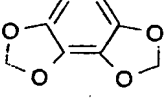
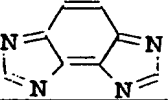
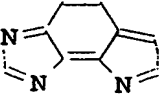
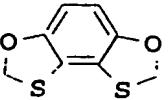
Table 1

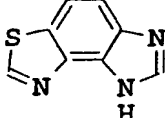
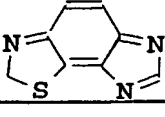
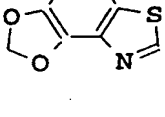
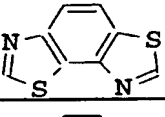
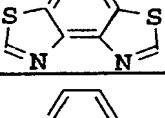
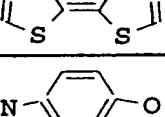
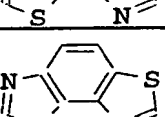
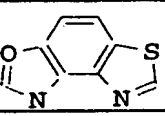
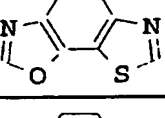
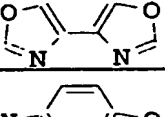
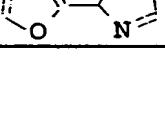

<u>Structure</u>	<u>Name</u>	<u>CA Reg. No.</u>	<u>Reference</u>
	benzo (2,1-b:3,4-b) dithiophene	211-53-0	Sturaro et al., <u>Heterocycl. Chem.</u> 27:1867, 1990
	benzo (2,1-b:3,4-b) difuran	211-47-2	Rene et al., <u>Eur. J. Med. Chem.-Chim. Ther.</u> 13:435, 1978
	thieno (3,2-g) benzofuran	438-31-9	Cagniant and Kirsch, <u>Hebd. Seances Acad. Sci. C.</u> 282:465, 1976
	2H-furo (3,2-g) indole	103671-62-1	Lawrence Jr., <u>Eur. Pat. Appl.</u> EP 173,520, 1986
	2H-benzo (2,1-b:3,4-b') dipyrrole	112149-08-3	Berlin et al., <u>J. Chem. Soc. Chem. Commun.</u> (15):1176, 1987

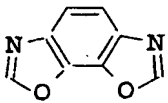
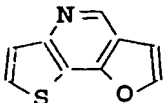
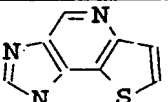
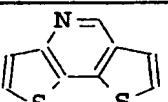
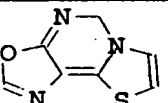
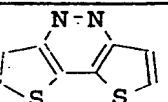
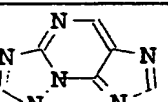
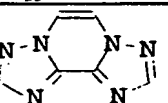
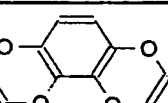
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	furo (3,2-h) quinoline	234-28-6	
	2,2'-bipyridyl	366-18-7	

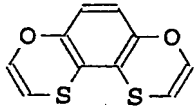
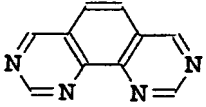
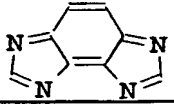
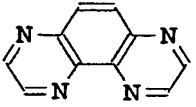
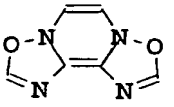
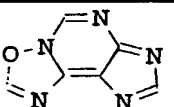
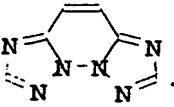
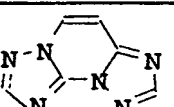
In structures I through VII above, further ring substitutions with heteroatoms are permitted. Preferably, such heteroatoms are selected from nitrogen, oxygen, sulfur, and phosphorus. For example, the compounds listed in Table 2 illustrate further representative multi-dentate ligands of the present invention having additional ring substitutions. As with Table 1, Table 2 identifies the structure of the representative multi-dentate ligands, lists the corresponding chemical name, identifies the CA Reg. No., and provides a corresponding reference (if available) describing the synthesis and/or chemistry of the identified multi-dentate ligand.

Table 2

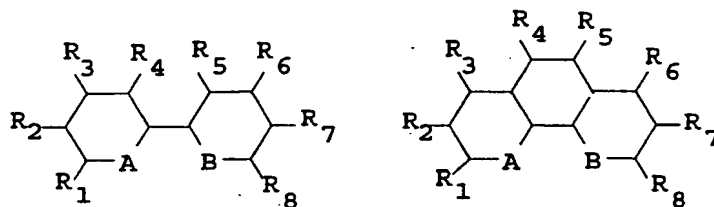
Structure	Name	CA Reg. No.	Reference
	furano (3,2-g) benzoxazole	25885-39-6	
	furano (2,3-e) benzoxazole	66037-80-1	Turin et al., Fr. Demande 2,338,041, 1977
	thieno (3,2-g) benzoxazole	58188-85-5	Iddon et al., <u>J. Chem. Soc., Perkin Trans. I</u> 17:1686, 1975
	thieno (3,2-g) benzothiazole	72121-58-5	
	thieno (2,3-e) benzothiazole	211-36-9	
	benzo (1,2-d:3,4- d') bis (1,3) dioxide	211-50-7	Dallacker and Weiner, <u>Justus Liebig's Ann. Chem.</u> 725:99, 1969
	benzo (1,2-d:3,4- d') diimidazole	211-10-9	
	pyrrolo(2,3-e) benzimidazole	53068-46-5	Chetverikov et al., U.S.S.R. 425,906, 1974
	benzo (2,1-d:3,4- d') bis (1,3) oxathiole	211-54-1	

	2H-imidazo (4,5-e) benzothiazole	42341-40-2	
	2H-imidazo (4,5-g) benzothiazole	211-23-4	
	1,3-dioxolo (4,5-e) benzothiazole	77482-58-7	Foerster et al., Ger. Offen. 2,903,966, 1980
	benzo (1,2-d:3,4-d') bisthiazole	211-37-0	
	benzo (2,1-d:3,4-d') bisthiazole	23147-19-5	
	benzo (1,2-d:4,3-d') bisthiazole	10558-80-2	Grandolini et al., Ann. Chim. 58:91, 1968
	thiazolo(5,4-e) benzoxazole	211-35-8	
	thiazolo (5,4-g) benzoxazole	51273-21-3	
	thiazolo (4,5-e) benzoxazole	315-47-9	
	thiazolo (4,5-f) benzoxazole	67239-73-0	Fridman et al., Ukr. Khim. Zh. 44:399, 1978
	benzo (2,1-d:3,4-d') bisoxazole	211-19-8	
	benzo (1,2-d:3,4-d') bisoxazole	211-20-1	

	benzo (1,2-d:4,3-d') bisoxazole	54935-19-2	Barker et al., <u>J. Chem. Res. Synop.</u> (9):328, 1986
	furo (2,3-d) thieno (3,2-b) pyridine	110665-19-5	
	1H-imidazo (4,5-d) thieno (3,2-b)-pyridine	111163-54-3	Takada et al., Eur. Pat. Appl. EP 223,420, 1987
	dithieno (3,2-b:2',3'-d) pyridine	40826-38-8	Yang et al., <u>Synthesis</u> 2:130, 1989; Heeres et al., <u>Syn. Commun.</u> 2:365, 1972
	5H-oxazolo (4,5-e) thiazolo (3,2-c) pyrimidine	211-46-1	
	dithieno (3,2-c:2',3'-e) pyridazine	51974-92-6	Nonciaux et al., Bull. Soc. Chim. Fr. 12 Pt 2, 3318, 1973
	1H-(1,2,4) triazolo (5,1-b) purine	387-96-2	
	bis (1,2,4) triazolo (1,5-d:5',1'-c) pyrazine	55366-22-8	Vercek et al., <u>Tetrahedron Lett.</u> (51/52):4539, 1974
	benzo (2,1-b:3,4-b') dipyran	231-29-8	<u>Monatsch</u> 80:743, 1949

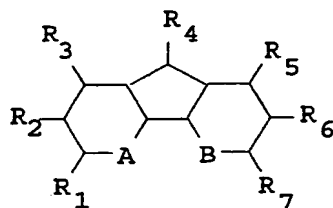
	benzo (1,2-b:4,3-b') bis (1,4)-oxathiin	231-34-5	
	benzo (1,2-e:3,4-e') dipyrazine		
	benzo (1,2-d:3,4-d') diimidazole	211-10-9	
	pyrazino (2,3-f) quinoxaline	231-23-2	Shim et al., <u>Synthesis</u> 2:116, 1980; Nasielski-Hinkins et al., <u>J. Chem. Soc. Perkin Trans.</u> 1:1229, 1975
	bis (1,2,4) oxadiazolo (2,3-d:3',2'-c) pyrazine	74382-83-5	
	(1,2,4)-oxadiazolo (3,2-i) purine	56248-95-4	Miura et al., <u>Chem. Pharm. Bull.</u> 23:464, 1975
	bis (1,2,4) triazolo (1,5-b:5',1'-f) pyridazine	51519-32-5	Polanc et al., <u>J. Org. Chem.</u> 39:2143, 1974
	bis (1,2,4) triazolo (1,5-d:1',5'-c) pyrimidine	76044-62-7	Brown and Shinozuka, <u>Aust. J. Chem.</u> 33:1147, 1980

General structures I through VII identified above may possess further chemical moieties covalently attached to the structural backbone, as illustrated below:

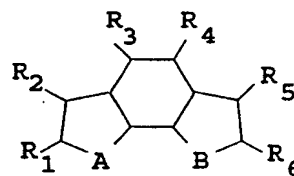


Ib

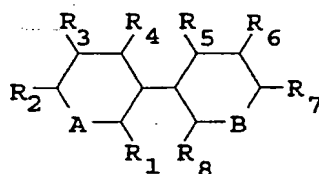
IIb



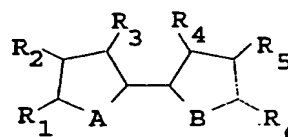
IIIb



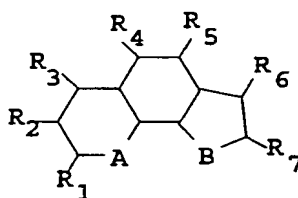
IVb



Vb



VIb



VIIb

wherein  $R_1$  through  $R_8$  are the same or different, and are selected from the following chemical moieties: -H, -OH, -X, -OX, -COOH, -COOX, -CHO, -CXO, -F, -Cl, -Br, -I, -CN,

-NH<sub>2</sub>, -NHX, -NX<sub>2</sub>, -PX<sub>2</sub>, -SO<sub>3</sub>H, -SO<sub>3</sub>Na, -SO<sub>3</sub>K, -SO<sub>3</sub>X, -PO<sub>3</sub>H, -OPO<sub>3</sub>H, -PO<sub>3</sub>X, -OPO<sub>3</sub>X and -NO<sub>2</sub>. As used herein, "X" represents and an alkyl moiety or an aryl moiety. An "alkyl moiety" is a straight chain or branched, cyclic or noncyclic, saturated or unsaturated, substituted or unsubstituted carbon chain containing from 1-20 carbon atoms; and an "aryl moiety" is a straight chain or branched, cyclic or noncyclic, saturated or unsaturated, substituted or unsubstituted carbon chain containing at least one substituted or unsubstituted aromatic moiety and containing from 6-20 carbon atoms. Such chemical moieties may also be covalently attached to the ring fusion atoms. Representative examples of the chemical moieties of this invention include, but are not limited to, the moieties identified in Table 3 below.

Table 3

-H	-CH <sub>3</sub>	-CH <sub>2</sub> Br
-CH <sub>2</sub> OH	-CH <sub>2</sub> Cl	-CBr <sub>3</sub>
-CH <sub>2</sub> C <sub>6</sub> H <sub>5</sub>	-C <sub>6</sub> H <sub>5</sub>	-(CH <sub>2</sub> ) <sub>1-12</sub> CH <sub>3</sub>
-Cl	-CHO	-COOH
-COOMe	-CH=NOH	-CH <sub>2</sub> NH <sub>2</sub>
-CH <sub>2</sub> C≡CH	-CH=CH <sub>2</sub>	-P(C <sub>6</sub> H <sub>5</sub> ) <sub>2</sub>
-CH <sub>2</sub> CH(CO <sub>2</sub> H) <sub>2</sub>	-CON(CH <sub>2</sub> COOH) <sub>2</sub>	-CH <sub>2</sub> N(CH <sub>2</sub> COOH) <sub>2</sub>
$\begin{array}{c} \text{-CH}_2 \\   \\ \text{N} \\   \\ \text{CH}_2\text{OH} \\   \\ \text{H} \end{array}$	$\begin{array}{c} \text{CH}_3 \\   \\ \text{-N-CH-CH-C}_6\text{H}_5 \\   \quad   \\ \text{CH}_3 \quad \text{OH} \end{array}$	$\begin{array}{c} \text{-CH}_2\text{N(CH}_2\text{)}_{11}\text{CH}_3 \\   \\ \text{CH}_3 \end{array}$
-Ph-SO <sub>3</sub> Na		

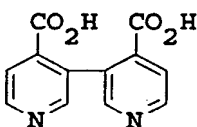
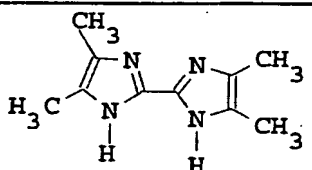
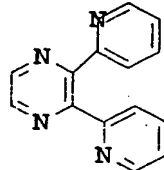

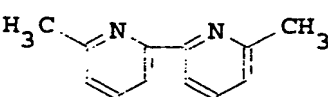
Representative examples of the multi-dentate ligands possessing further chemical moieties covalently attached to the structural backbone of structures I through VII are presented in Table 4. In particular, Table 4 identifies the structure of the representative multi-dentate ligands, lists the corresponding chemical name, identifies the CA



Reg. No., and provides a corresponding reference (if available) describing the synthesis and/or chemistry of the multi-dentate ligand.

5

Table 4

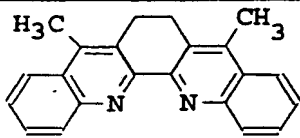
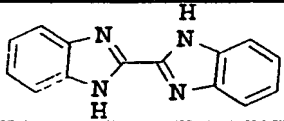
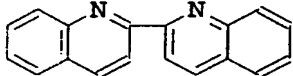
Structure	Name	CA Reg. No.	Reference
	2,2'-bipyridine-4,4'-dicarboxylic acid	6813-38-3	
	2,2'-bis (4,5-dimethylimidazole)	69286-06-2	J. Organomet. Chem. 307:39, 1986
	2,3-bis (2-pyridyl)pyrazine	25005-96-3	(Aldrich: 28,164-16)
	5,5'-dimethyl-2,2'-bithiophene	16303-58-5	
	6,6'-dimethyl-2,2'-dipyridine	4411-80-7	Kauffmann et al., Chem. Ber. 109:3864, 1976

10 The chemical moieties covalently attached to the structural backbone may be joined to yield an aromatic or nonaromatic cyclic chemical moiety. Representative examples of such cyclic chemical moieties are set forth in Table 5, which identifies the structure of the representative multi-dentate ligands, lists the

corresponding chemical name, identifies the CA Reg. No., and provides a corresponding reference (if available) describing the synthesis and/or chemistry of the multi-dentate ligand.

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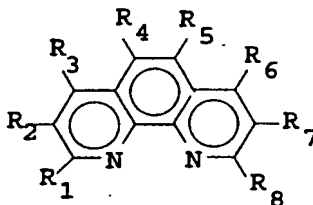
Table 5

<u>Structure</u>	<u>Name</u>	<u>CA Reg. No.</u>	<u>Reference</u>
	6,7-dihydro- 5,8-dimethyl dibenzo (b) (1,10) phenanthroline	5298-71-5	
	bibenzimidazole	123067-51-6	
	2,2'- bisquinoline	119-91-5	(Aldrich: B3,540-7)

The synthesis of representative examples of the multi-dentate ligands of this invention are disclosed in Table 6 and Table 7 below. Specifically, in these tables the structure of the multi-dentate ligands are identified along with their CA Reg. No. and one or more references disclosing their synthesis and/or chemistry.

15

Table 6  
Synthesis of Representative Copper(I) Complexes  
Having the Structure:

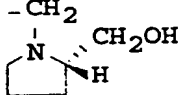
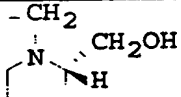
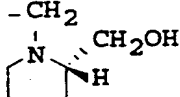
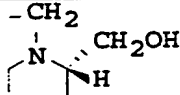
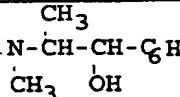


5

(R<sub>2</sub> through R<sub>7</sub> = hydrogen, unless indicated)

R <sub>1</sub>	R <sub>8</sub>	CA Reg. No.	Reference
-CH <sub>3</sub>	-CH <sub>3</sub>	484-11-7	O'Reilly et al., <i>Aust. J. Chem.</i> 13:145, 1960
-CH <sub>2</sub> Br	-CH <sub>2</sub> Br	78831-37-5	Weijen et al., <i>J. Org. Chem.</i> 57:7258, 1992; Jukkala et al., <i>Helv. Chim. Acta.</i> 75:1621, 1992; Chandler et al., <i>J. Heterocycl. Chem.</i> 18:599, 1981
-CH <sub>2</sub> Br	-CH <sub>2</sub> OH	142470-16-4	Weijen et al., <i>J. Org. Chem.</i> 57:7258, 1992
-CBr <sub>3</sub>	-CBr <sub>3</sub>		Chandler et al., <i>J. Heterocycl. Chem.</i> 18:599, 1981
-CH <sub>2</sub> Cl	-CH <sub>2</sub> Cl		Newcome et al., <i>J. Org. Chem.</i> 50:3807, 1985; Newcome et al., <i>J. Org. Chem.</i> 48:5112, 1983
-CCl <sub>3</sub>	-CCl <sub>3</sub>		Chandler et al., <i>J. Heterocycl. Chem.</i> 18:599, 1981; Newcome et al., <i>J. Org. Chem.</i> 48:5112, 1983
-CN	-CN	57709-63-4	Chandler et al., <i>J. Heterocycl. Chem.</i> 18:599, 1981; Sjoegren et al., <i>Organometallics</i> 11:3954, 1992

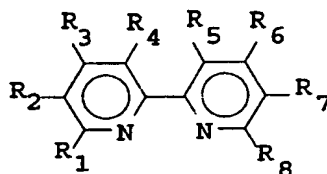
-CH <sub>2</sub> C <sub>6</sub> H <sub>5</sub>	-CH <sub>2</sub> C <sub>6</sub> H <sub>5</sub>	223-20-1	Sjoegren et al., <u>Organometallics</u> 11:3954, 1992
-(CH <sub>2</sub> ) <sub>11</sub> CH <sub>3</sub>	-(CH <sub>2</sub> ) <sub>11</sub> CH <sub>3</sub>		Menger et al., <u>J. Am. Chem. Soc.</u> 113:4017, 1991
-(CH <sub>2</sub> ) <sub>3</sub> CH <sub>3</sub>  (R <sub>3</sub> =R <sub>6</sub> =H, Ph)	-(CH <sub>2</sub> ) <sub>3</sub> CH <sub>3</sub>	85575-93-5P	Sugihara et al., JP 02096578 A2, <u>Jpn. Kokai Tokkyo Koho</u> 113(15):132159v
-(CH <sub>2</sub> ) <sub>3</sub> CH <sub>3</sub>  (R <sub>4</sub> =R <sub>5</sub> =-CH <sub>3</sub> )	-(CH <sub>2</sub> ) <sub>3</sub> CH <sub>3</sub>		Delton et al., EP 339973 A1, <u>Eur. Pat. Appl.</u> 112(21):19835p, 1989
-Cl	-Cl	29176-55-4	Sjoegren et al., <u>Organometallics</u> 11:3954, 1992; Delton et al., EP 339973 A1, <u>Eur. Pat. Appl.</u> 112(21):19835p, 1989
-CH <sub>2</sub> OH	-CH <sub>2</sub> OH	78831-36-4	Chandler et al., <u>J. Heterocycl. Chem.</u> 18:599, 1981; Delton et al., EP 339973 A1, <u>Eur. Pat. Appl.</u> 112(21):19835p, 1989; Newcome et al., <u>J. Org. Chem.</u> 48:5112, 1983
-CHO	-CHO	57709-62-3	Ziessel, <u>Tetrahedron Lett.</u> 30:463, 1989; Toner, EP 288256 A2, <u>Eur. Pat. Appl.</u> 111(15):130322c; Bell et al., <u>J. Inclusion Phenom.</u> 5:149, 1987
-COOH	-COOH		Chandler et al., <u>J. Heterocycl. Chem.</u> 18:599, 1981
-COOMe	-COOMe		Chandler et al., <u>J. Heterocycl. Chem.</u> 18:599, 1981; Newcome et al., <u>J. Org. Chem.</u> 48:5112, 1983

-CH=NOH	-CH=NOH		Chandler et al., <i>J. Heterocycl. Chem.</i> 18:599, 1981
-CH <sub>2</sub> NH <sub>2</sub>	-CH <sub>2</sub> NH <sub>2</sub>		Chandler et al., <i>J. Heterocycl. Chem.</i> 18:599, 1981
-CHO	-H	33795-37-8	Toner, EP 288256 A2, <i>Eur. Pat. Appl.</i> 111(15):130322c
-COOH	-H	1891-17-4	Toner, EP 288256 A2, <i>Eur. Pat. Appl.</i> 111(15):130322c
-CH <sub>2</sub> C≡CH	-CH <sub>2</sub> C≡CH		Sjoegren et al., <i>Organometallics</i> 11:3954, 1992
-C <sub>6</sub> H <sub>5</sub>	-C <sub>6</sub> H <sub>5</sub>		Dietrich-Buchecker et al., <i>Tetrahedron Lett.</i> 23:5291, 1982
-Cl	-CH <sub>3</sub>		Newcome et al., <i>J. Org. Chem.</i> 54:1766, 1989
-CH=CH <sub>2</sub>	-CH=CH <sub>2</sub>		Newcome et al., <i>J. Org. Chem.</i> 50:3807, 1985
-P(C <sub>6</sub> H <sub>5</sub> ) <sub>3</sub>	-P(C <sub>6</sub> H <sub>5</sub> ) <sub>3</sub>		Ziessel, <i>Tetrahedron Lett.</i> 30:463, 1989
CH <sub>2</sub> CH(CO <sub>2</sub> H) <sub>2</sub>	-CH <sub>2</sub> CH(CO <sub>2</sub> H) <sub>2</sub>		Newcome et al., <i>Inorg. Chem.</i> 24:811, 1985
-CH <sub>2</sub> N(CH <sub>2</sub> ) <sub>11</sub> CH CH <sub>3</sub>			Weijen et al., <i>J. Org. Chem.</i> 57:7258, 1992
			Weijen et al., <i>J. Org. Chem.</i> 57:7258, 1992
-CH <sub>2</sub> OH			Weijen et al., <i>J. Org. Chem.</i> 57:7258, 1992
-CH <sub>2</sub> N(CH <sub>2</sub> ) <sub>11</sub> CH CH <sub>3</sub>			Weijen et al., <i>J. Org. Chem.</i> 57:7258, 1992
-CH <sub>2</sub> N(CH <sub>2</sub> COOH) <sub>2</sub>	-CH <sub>2</sub> N(CH <sub>2</sub> COOH) <sub>2</sub>		Mukkala et al., <i>Helv. Chim. Acta</i> 75:1621, 1992; Toner, EP 288256 A2, <i>Eur. Pat. Appl.</i> 111(15):130322c

-CON(CH <sub>2</sub> COOH) <sub>2</sub>	-CON(CH <sub>2</sub> COOH) <sub>2</sub>		Toner, EP 288256 A2, <u>Eur. Pat. Appl.</u> 111(15):130322c.
-CH <sub>3</sub> (R <sub>3</sub> =R <sub>6</sub> = -Ph-SO <sub>3</sub> Na	-CH <sub>3</sub>	52698-84-7	Blair et al, <u>Talanta</u> 2:163, 1961

Table 7

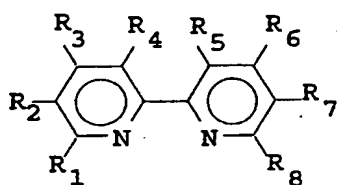
Synthesis of Representative Copper(I) Complexes  
Having the Structure:



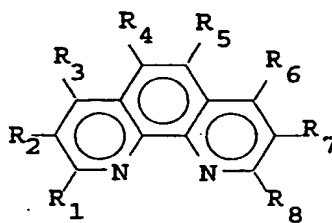
(R<sub>2</sub> through R<sub>7</sub> = hydrogen, unless indicated)

R1	R8	CA Reg. No.	Reference
-CN	-CN	4411-83-0	Sjoegren et al., <u>Organometallics</u> 11:3954, 1992
-CH <sub>2</sub> Cl	-CH <sub>2</sub> Cl	74065-64-8	Bell et al., <u>J. Inclusion Phenom.</u> 5:149, 1987
-CHO	-CHO		Newkome et al., <u>J. Org. Chem.</u> 50:3807, 1985
-CH=CH <sub>2</sub>	-CH=CH <sub>2</sub>		Newkome et al., <u>J. Org. Chem.</u> 50:3807, 1985
(R <sub>1</sub> and R <sub>2</sub> = benzo moiety)	(R <sub>7</sub> and R <sub>8</sub> = benzo moiety)	119-91-5	(Aldrich: B3,540-7)

In one embodiment of this invention, the multi-dentate ligands are selected from the following structures:



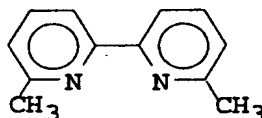
Ic



IIc

wherein  $R_1$  through  $R_8$  are the same or different, and are selected from hydrogen, an alkyl moiety and an aryl moiety.

In a preferred embodiment, the multi-dentate ligand is 6,6'-dimethyl-2,2'-dipyridine having structure Id:

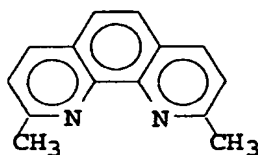


Id

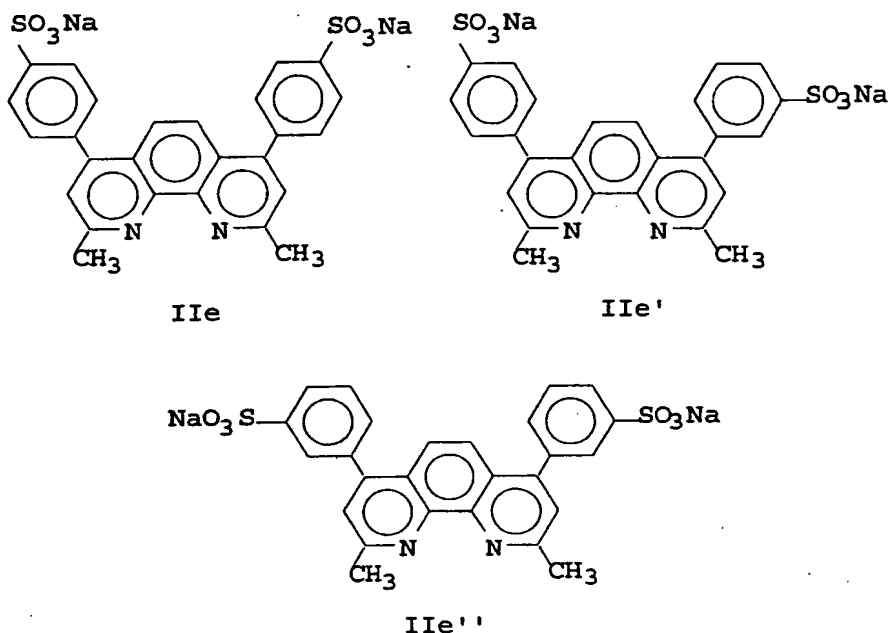
10

In a further preferred embodiment, the multi-dentate ligand is neocuproine (2,9-dimethyl-1,10-phenanthroline) having structure IIId, or is bathocuproine disulfonic acid ("BCDS") having one of the isomeric structures IIe, IIe' or IIe'':

15



IIId



5

Unless otherwise indicated, BCDS refers to a physical mixture of the above isomers (i.e., IIe, IIe' and IIe''). Typically, the ratio of the various isomers (i.e.,  
 10 IIe:IIe':IIe'') vary depending upon the commercial source of BCDS as follows: Aldrich Chemical Co., Inc. (Milwaukee, Wisconsin) 9.1:38.6:41.2; Spectrum Chemical Manufacturing Corp. (Gardena, California) 8.5:39.7:45.2; GFS Chemicals (Columbus, Ohio) 8.4:38.5:45.3; Janssen Pharmaceutica  
 15 (subsidiary of Johnson & Johnson) (Beerse, Belgium) 4.6-8.7:36.4-39.4:44.4-55.9.

As discussed above, stable copper(I) complexes of this invention may be made by contacting a multi-dentate ligand with a copper(I) source. The multi-dentate ligands  
 20 may be obtained from commercial sources, or may be synthesized by known organic synthesis techniques from commercially available reagents. Preferably, water soluble multi-dentate ligands are complexed with the copper(I) in aqueous solution, employing CuCl, Cu<sub>2</sub>O or  
 25 CuCN as the copper(I) source. The resulting copper(I)



complex may then be recovered by evaporation of solvent to yield the copper(I) complex. Alternatively, if the multi-dentate ligand is not readily soluble in water, copper(I) complexes may be formed by the above procedure employing a  
5 suitable non-aqueous (e.g., organic) solvent.

In the practice of this invention, the ratio of the multi-dentate ligand to copper(I) may be any ratio which results in a stable copper(I) complex. Preferably, the ligand to copper ratio is at least 1:1. In a more  
10 preferred embodiment, the ligand to copper ratio ranges from 1:1 to 3:1 (including 2:1). Such copper(I) complexes may be made by the procedures identified in the preceding paragraph by reacting the appropriate molar ratios of the multi-dentate ligand and the copper(I) ion source.

15 Although not intending to be limited by the following theory, it is believed that copper(I) has enhanced biological activity over copper(II) in certain biological events. For example, it is believed that copper(I) may be an important intermediate for copper metabolism, including  
20 copper uptake and/or transfer, as well as cellular delivery. Thus, the reduction of copper(II) to copper(I) is bypassed by direct delivery of copper(I). Furthermore, the stable copper(I) complexes of this invention are suitable for systemic delivery to warm blooded animals,  
25 and may provide a sustained release of copper to the animal.

The stable copper(I) complexes of this invention possess utility as therapeutic substances, including utility as anti-oxidative and anti-inflammatory agents  
30 generally and, more specifically, as wound healing agents. The copper(I) complexes of this invention also possess activity as hair growth agents, lipid modulation agents, signal transduction modulating agents, and anti-viral agents. For purpose of clarity, the various biological  
35 activities of the stable copper(I) complexes of this invention are addressed individually below.

Highly reactive oxygen species such as the superoxide anion ( $O_2^{\bullet-}$ ), hydrogen peroxide ( $H_2O_2$ ), hydroxyl radical ( $HO^\bullet$ ), and lipid peroxides (LOOH) are involved in a number of human diseases. For example, such oxygen species have been implicated in autoimmune diseases, arthritis, tissue damage caused by environmental pollutants, cigarette smoke and drugs, tissue injury during, for example, surgery and transplantation, as well as a variety of other conditions (see, e.g., Halliwell, B., Fed. Amer. Soc. Exp. Biol. 1:358-364, 1987). Reactive oxygen species are also generated during the response to injury by phagocytic cells. One of the early events in the wound healing response is the cleansing and sterilization of the wound by neutrophils and macrophages. A mechanism for this sterilization is the generation of the superoxide anion and hydrogen peroxide, and generally results in an inflammatory response. Moreover, superoxide anion and hydrogen peroxide will, in the presence of iron or other redox active transition metal complexes, generate the hydroxyl radical. The hydroxyl radical is a potent oxidant which initiates the free radical oxidation of fatty acids, as well as the oxidative degradation of other biomolecules. For example, an important area in which reactive oxygen species cause tissue damage is in post-injury damage to the brain and spinal chord, and in reperfusion injury to ischemic tissue following surgery and transplantation (such as heart surgery and/or transplantation). A sudden inrush of oxygenated blood and activated phagocytic cells leads to superoxide anion and hydrogen peroxide formation. These species do direct damage to tissue, and also react with iron (as discussed above) to generated the very reactive hydroxyl radical.

The stable copper(I) complexes of this invention generally serve as anti-oxidative agents which prevent or limit the oxidative damage caused by reactive oxygen species, and further serve as anti-inflammatory agents by

reducing the inflammatory response associated with such reactive oxygen species. More specifically, the copper(I) complexes of the present invention are useful in the enhancement and/or restoration of the defense of warm-blooded animals to oxidative or inflammatory damage caused by the highly reactive oxygen species, and may be used in pharmaceutical preparations to inhibit oxidative and inflammatory processes which lead to tissue damage. Moreover, the stable copper(I) complexes of this invention accelerate the wound healing process by "detoxifying" tissue damage by the highly reactive oxygen species.

In addition to highly reactive oxygen species, macrophages and neutrophils induce or continue an inflammatory response through the generation of certain lipid mediators of inflammation (e.g., leukotrienes and prostaglandins). The involvement of such mediators in inflammatory bowel disease (IBD) and related chronic inflammatory conditions, such as arthritis, is evidenced by a strong correlation between disease progression and the levels and presence of leukotrienes and prostaglandins in the circulation and effected tissue. Prostaglandins enhance vasodilation and edema formation, while leukotrienes are potent chemoattractive agents for leukocytes, especially neutrophils, and stimulate degranulation and the release of damaging lysosomal enzymes and superoxide production.

The distribution of the two major pathways leading either to prostaglandins or to leukotrienes varies according to cell type. While most cells possess the cyclooxygenase pathway, the 5-lipoxygenase pathway leading to the leukotrienes is less widely distributed and is prominent in inflammatory cells, such as neutrophils, macrophages, monocytes and mast cells. The general scheme for lipid mediator synthesis is illustrated in Figure 3.

The stable copper(I) complexes of this invention inhibit the formation of prostaglandins and/or

leukotrienes by inhibiting the enzymes involved in their formation. Referring to Figure 3, the stable copper(I) complexes are effective inhibitors of both cyclooxygenase-1 and cyclooxygenase-2, thereby inhibiting the formation of prostaglandins. Similarly, the stable copper(I) complexes are effective inhibitors of 5-lipoxygenase and leukotriene C<sub>4</sub> (LCT<sub>4</sub>) synthetase, thereby inhibiting the formation of leukotrienes.

In addition, proteolysis of various cellular targets by elastase (a neutrophil-released serine protease) at the site of inflammation has been implicated in a number of pathologic conditions, including emphysema, rheumatoid arthritis, and psoriasis. Thus, inhibitors of elastase may be used to treat, prevent or limit the breakdown of normal tissue at the site of inflammation, and the stable copper(I) complexes of this invention are effective inhibitors of elastase.

The stable copper(I) complexes of this invention may also be used in the regulation and/or modulation of lipid metabolism in general. For example, hypercholesterolemia and hyperlipidemia are common and serious health problems which are treatable with the stable copper(I) complexes of this invention.

Hypercholesterolemia has been observed in marginal and severely copper-deficient rats, as well as other animals, including humans (Lei, "Plasma Cholesterol Response in Copper Deficiency," Role of Copper in Lipid Metabolism, ed. Lei, CRC Press, pages 1-24, 1990). Elevation in serum cholesterol level has been linked to increases in the activity of hepatic 3-hydroxy-3-methylglutaryl coenzyme A reductase (HMG CoA reductase, E.C.1.1.1.34) and glutathione levels (Bunce, "Hypercholesterolemia of Copper Deficiency is Linked to Glutathione Metabolism and Regulation of HMG CoA Reductase," Nutr. Rev. 51: 305-307, 1993; Kim et al., "Inhibition of Elevated Hepatic Glutathione Abolishes

Copper Deficiency Cholesterolemia," FASEB J. 6: 2467-2471, 1992).

Similar increases in the synthesis and level of other hepatic lipids (fatty acids, triacylglycerols and phospholipids) have been observed in copper deficient rats (al-Othman et al., "Copper Deficiency Increases In Vivo Hepatic Synthesis of Fatty Acids, Triacylglycerols, and Phospholipids," Proc. Soc. Exp. Biol. Med. 204(1): 97-103, 1993) and treatment with a copper(II) complex has been shown to lower the activity of liver enzymes involved in lipid metabolism, including acetyl CoA synthetase in vivo (Hall et al., "Hypolipidemic Activity of Tetetrakis-mu-(trimethylamine-boranecarboxylato)-bis(trimethylamine-carboxylborane)-dicopper(II) in Rodents and its Effect on Lipid Metabolism," J. Pharm. Sci. 73(7): 973-977, 1984). Conversely, it has been reported that treatment by injection of copper(II) increased serum cholesterol concentrations in rats, possibly by increasing the activity of the HMG CoA reductase (Tanaka et al., "Effect of Cupric Ions on Serum and Liver Cholesterol Metabolism," Lipids 22: 1016-1019, 1987). Accordingly, it is believed that copper may be an important factor in the regulation of lipid levels.

Acetyl CoA synthetase catalyzes the formation of acetyl CoA from acetate. As illustrated in Figure 4, acetyl CoA can be further metabolized along many different pathways leading primarily to the formation of cholesterol and fatty acids or energy production. Agents which inhibit this enzyme influence the biosynthesis of various lipids. HMG-CoA reductase (3-hydroxy-3-methylglutaryl coenzyme A reductase) is located biochemically later in the lipid synthesis scheme and converts HMG-CoA to mevalonic acid, and is the rate limiting reaction in cholesterol biosynthesis (see Figure 4). Stable copper(I) compounds of this invention inhibit certain key enzymes involved in the formation of lipids, and thus serve as

lipid modulating or regulating agents. (The ability of stable copper(I) complexes to inhibit enzymes in the formation of lipids is disclosed in further detail in Examples 12-13.)

5       The stable copper(I) complexes of this invention may also serve as modulating agents of signal transduction in cells. Most intracellular signaling processes are regulated by reversible phosphorylation of specific proteins by kinases. Breakdown of phosphatidylinositol  
10 leads to the formation of diacylglycerol and inositol triphosphate, the former acting synergistically with calcium to activate Protein Kinase C (PKC), resulting in translocation of the enzyme from cytosol to the membrane. Phosphorylation of proteins by PKC has been implicated as  
15 a pivotal regulatory element in signal transduction, cellular regulation and tumor promotion. Inhibitors of PKC, as well as other protein kinases, have the potential to block proliferative signaling in tumor induction, atherosclerosis and immune modulation.

20       Examples of factors which stimulate the G-protein linked phospholipase C breakdown of phosphatidylinositol include angiotensin II, bradykinin, endothelin, f-Met-Leu-Phe, and vasopressin. These protein kinase C enzymes are also directly activated by tumor promoters such as phorbol  
25 esters. Examples of Receptor linked tyrosine kinases include Epidermal Growth Factor, Nerve Growth Factor, and Platelet Derived Growth Factor. Examples of cytoplasmic tyrosine kinase activators include cytokines such as Interleukin 2, Interleukin 3, and Interleukin 5. These  
30 factors bind to specific lymphocyte receptors which activate the cytoplasmic tyrosine kinase.

      The action of PKC and protein tyrosine kinase action is illustrated in Figure 5. The stable copper(I) complexes of this invention serve as signal transduction modulating  
35 agents by inhibiting one or more enzymes involved in

intracellular signal transduction, including PKC and protein tyrosine kinases.

When administered to an animal to treat the conditions discussed above, the stable copper(I) complexes may first be combined with one or more suitable carriers or diluents to yield a pharmaceutical preparation suitable for topical, oral or parenteral application. Such diluents or carriers, however, should not interact with the stable copper(I) complex to significantly reduce the effectiveness thereof, or oxidize copper(I). Effective administration will preferably deliver a dosage of approximately 0.01 to 100 mg of the stable copper(I) complex per kg of body weight.

Methods for encapsulating compositions (such as in a coating of hard gelatin) for oral administration are well known in the art (see, e.g., Baker, Richard, Controlled Release of Biological Active Agents, John Wiley and Sons, 1986) (incorporated herein by reference). Suitable carriers for parenteral application (such as intravenous, subcutaneous or intramuscular injection) include sterile water, physiological saline, bacteriostatic saline (saline containing 0.9 mg/ml benzyl alcohol) and phosphate-buffered saline. The stable copper(I) complexes may be topically applied in the form of liquids, containing pharmaceutically acceptable diluents (such as saline and sterile water) or may be applied as lotions, creams or gels, containing additional ingredients to impart the desired texture, consistency, viscosity and appearance. Such additional ingredients are familiar to those skilled in the art and include emulsifying agents such as non-ionic ethoxylated and nonethoxylated surfactants, fatty alcohols, fatty acids, organic or inorganic bases, preserving agents, wax esters, steroid alcohols, triglyceride esters, phospholipids such as lecithin and cephalin, polyhydric alcohol esters, fatty alcohol esters, hydrophilic lanolin derivatives, hydrophilic beeswax

derivatives, hydrocarbon oils such as palm oil, coconut oil, mineral oil, cocoa butter waxes, silicon oils, pH balancers and cellulose derivatives.

Topical administration may be accomplished by  
5 applying an amount of the preparation directly to the desired area, such as a wound or an inflamed area. The required dosage will vary according to the particular condition to be treated, the severity of the condition, and the duration of the treatment. Preferably, when the  
10 stable copper(I) complex is topically applied in the form of a lotion, cream or gel, the preparation may contain about 1% to about 20% of a penetration enhancing agent. Examples of penetration enhancing agents include dimethylsulfoxide (DMSO), urea and eucalyptol. In the  
15 case of a liquid preparations for topical application, the concentration of penetration enhancing agent (such as DMSO) may comprise about 30% to about 80% of the preparation.

In addition to the activity discussed above, the  
20 stable copper(I) complexes of this invention also possess utility as hair growth agents. Hair loss is a common affliction of humans, the most common being "alopecia" where males lose scalp hair as they get older (also called "male pattern baldness"). Other hair loss afflictions  
25 include alopecia areata (AA), female pattern baldness and secondary alopecia (e.g., hair loss associated with chemotherapy and/or radiation treatment). The stable copper(I) complexes of this invention are particularly useful in stimulating hair growth associated with any hair  
30 loss affliction, including the specific afflictions identified above.

Hair is normally divided into two types, "terminal" and "vellus" hairs. Terminal hair is coarse, pigmented hair which arises from follicles which are developed deep  
35 within the dermis. Vellus hairs are typically thin, non-pigmented hairs which grow from hair follicles which are



smaller and located superficially in the dermis. As alopecia progresses, there is a change from terminal to vellus type hair. Other changes that contribute to alopecia are alterations in the growth cycle of hair.

5 Hair typically progresses through three cycles, anagen (active hair growth), catagen (transition phase), and telogen (resting phase during which the hair shaft is shed prior to new growth). As baldness progresses, there is a shift in the percentages of hair follicles in each phase,

10 with the majority shifting from anagen to telogen. The size of hair follicles is also known to decrease while the total number remains relatively constant.

As mentioned above, the stable copper(I) complexes of this invention have utility as stimulating agents for the growth of hair in warm-blooded animals. In one embodiment

15 of the present invention, the copper(I) complex may be administered intradermally in the area to be treated, along with a suitable vehicle, at a concentration of approximately 100-500 micrograms of copper(I) complex per

20 0.1 ml of vehicle. Suitable vehicles in this regard include saline, sterile water, and the like.

In another embodiment, the stable copper(I) complex may be topically applied in the form of a liquid, lotion, cream or gel by applying an effective amount of the topical preparation directly to the scalp. Any quantity

25 sufficient to stimulate the rate of hair growth is effective, and treatment may be repeated as often as the progress of hair growth indicates. Preferably, suitable topical hair growth preparations contain from about 0.1% to about 20% by weight of the stable copper(I) complex

30 (based on the total weight of the preparation).

Topical hair growth preparations of the present invention may contain about 0.5% to about 10% of an emulsifying or surface active agent. Non-ionic surface

35 active agents and ionic surface active agents may be used for the purposes of the present invention. Examples of

suitable non-ionic surface active agents are nonylphenoxypolyethoxy ethanol (Nonoxynol-9), polyoxyethylene oleyl ether (Brij-97), various polyoxyethylene ethers (Tritons), and block copolymers of ethylene oxide and propylene oxide of various molecular weights (Pluronic 68, for example). Acceptable preparations may also contain about 1% to about 10% of certain ionic surface active agents. These ionic surface active agents may be used in addition to or in place of, the non-ionic surface active agents. Examples of ionic surface active agents are sodium lauryl sulfate and similar compounds.

In addition to, or in place of, the emulsifying or surface active agent, topical hair growth preparations of this invention may contain about 1% to about 20% of a penetration enhancing agent. Examples of penetrating enhancing agents are DMSO and Urea. In the case of a liquid preparation to be applied topically, the concentration of a penetrating enhancing agent, such as DMSO, may comprise about 30% to about 80% of the topical preparation. The balance of the topical hair growth preparation may comprise an inert, physiologically acceptable carrier. Suitable carriers include, but are not limited to, water, physiological saline, bacteriostatic saline (saline containing 0.9 mg/ml benzyl alcohol), petrolatum based creams (e.g., USP hydrophilic ointments and similar creams, Unibase, Parke-Davis), various types of pharmaceutically acceptable gels, and short chain alcohols and glycols (e.g., ethyl alcohol and propylene glycol).

The following are examples of suitable hair growth preparations within the context of the present invention:

Preparation A:

35	Copper(I) Complex	10.0% (w/w)
	Hydroxy Ethyl Cellulose	3.0%

	Propylene Glycol	20.0%
	Nonoxynol-9	3.0%
	Sodium Lauryl Sulfate	2.0%
	Benzyl Alcohol	2.0%
5	0.2N Phosphate Buffer	60.0%
	<u>Preparation B:</u>	
	Copper(I) Complex	10.0% (w/w)
	Nonoxynol-9	3.0%
	Ethyl Alcohol	87.0%
10	<u>Preparation C:</u>	
	Copper(I) Complex	5.0% (w/v)
	Ethyl Alcohol	47.5%
	Isopropyl Alcohol	4.0%
	Propylene Glycol	20.0%
15	Laoneth-4	1.0%
	Water	22.5%
	<u>Preparation D:</u>	
	Copper(I) Complex	5.0% (w/v)
	Water	95.0%
20	<u>Preparation E:</u>	
	Copper(I) Complex	5.0% (w/v)
	Hydroxypropyl Cellulose	2.0%
	Glycerin	20.0%
	Nonoxynol-9	3.0%
25	Water	70.0%
	<u>Preparation F:</u>	
	Copper(I) Complex	1.0% (w/w)
	Nonoxynol-9	5.0%
	Unibase Cream	94.0%
30	<u>Preparation G:</u>	
	Copper(I) Complex	2.0% (w/w)
	Nonoxynol-9	3.0%
	Propylene Glycol	50.0%
	Ethanol	30.0%
35	Water	15.0%

The copper(I) complexes of the present invention also posses utility as anti-viral agents, and are particularly effective in the inhibition of the AIDS virus. Human acquired immunodeficiency syndrome or "AIDS" is a fatal disease for which there is presently no cure. The disease is believed to be caused by a virus known as the human immunodeficiency virus, commonly referred to as "HIV." The virus is transmitted by HIV-infected individuals through the exchange of bodily fluids. HIV infection results most commonly from sexual contact with an infected partner and the sharing among intravenous drug users of hypodermic syringes previously used by an infected individual. A pregnant HIV-infected mother may infect her unborn child by trans-placental transmission, and HIV-contaminated blood is a possible source of infection for individuals subject to blood transfusion.

HIV infection causes a suppression of the immune system. The immune suppression renders the infected individual vulnerable to a variety of opportunistic infections and conditions that are otherwise kept in balance by a healthy immune system. Fatalities result from HIV infection due to the inability of AIDS patients to respond to treatment of the opportunistic infections and conditions as a consequence of their compromised immune systems. Because the virus may often remain dormant, the manifestation of AIDS from HIV infection may take as long as ten years.

One approach to the treatment of AIDS has targeted the opportunistic infections or conditions which result from HIV infection. The treatment of such infections or conditions, however, is ultimately ineffective and, while prolonging the life of the infected individual, does not treat the underlying HIV infection. A second approach to the treatment of AIDS targets the cause of the disease itself. Because AIDS results from viral infection, it is believed that viral inactivation may ultimately provide a

cure. Materials which are capable of viral inactivation or inhibition are referred to herein as "antiviral agents."

To understand the mode of action of antiviral agents in the treatment of AIDS, an understanding of the process of HIV infection is necessary. HIV chronically infects specific immune cells known as T-helper cells, which are required for normal immune response. The HIV infected T-helper cells serve as hosts to the virus and facilitate the reproduction of the virus (the process of viral reproduction is commonly referred to as "replication"). After HIV infection, the infected host cell eventually dies, the replicated HIV virus is released, and the infection spreads to additional cells. This cycle continues unabated, depleting the population of T-helper cells and, in time, weakens the immune system to the onset of AIDS symptoms. Because T-helper cells are continuously produced by the body, the population of these cells may be reestablished in the absence of further HIV infection. Therefore, the progression of HIV infection (and the subsequent onset of AIDS) may be arrested by the prevention or inhibition of viral replication, and antiviral agents capable of inhibiting or preventing the replication of HIV should be effective in the treatment of AIDS.

At the genetic level, HIV replication requires the insertion of viral deoxyribonucleic acid ("DNA") into the genome of the host cell. The genome of the host cell consists of the cell's own DNA, and is responsible for the synthesis of materials essential to the cell's own function and proliferation. Once the viral DNA is inserted into the host genome, the host facilitates replication of HIV. The inserted viral DNA is an enzymatic product derived from viral ribonucleic acid ("RNA") and the action of an enzyme known as HIV reverse transcriptase. Inhibition of HIV reverse transcriptase

precludes the formation of viral DNA required for insertion into the genome of the host. Viral replication is prevented by the absence of viral DNA in the host cell genome. Antiviral agents which inhibit HIV reverse transcriptase are thus potential therapeutic drugs for treatment of AIDS.

Accordingly, in yet another embodiment of the present invention, antiviral agents are disclosed for inhibiting HIV replication, as well as methods relating to the administration thereof to an HIV-infected patient. The antiviral agents of this invention are the stable copper(I) complexes disclosed above, and the methods include administration of a therapeutically effective amount of a composition which includes a stable copper(I) complex in combination with a pharmaceutically acceptable carrier or diluent. Although not limited by the following theory, it is believed that the copper(I) complexes of this invention enhance transport of copper(I) into HIV infected cells which, in turn, inhibits or inactivates HIV protease and thus inhibits the replication of HIV. As used herein, the term "HIV" includes the various strains of the virus such as HIV-1 and HIV-2.

Administration of the stable copper(I) complexes of the present invention may be accomplished in any manner which will result in a systemic dose of a therapeutically effective amount of the copper(I) complex to an HIV-infected animal or patient (including human patients). For example, such administration may be by injection (intramuscular, intravenous, subcutaneous or intradermal), oral, nasal, or suppository applications. Typically, preparations of the present invention include stable copper(I) complexes in solution for various forms of injection, or in preparations which are formulated for the sustained release of the stable copper(I) complexes for oral, nasal, or suppository dosage application and generally include one or more inert, physiological

acceptable carriers. As used herein, the term "effective amount" means an amount of the stable copper(I) complex which inhibits HIV replication in the patient. Suitable dosages may range from approximately 0.01 to 100 mg of  
5 stable copper(I) complex per kg body weight.

The stable copper(I) complexes of this invention may be screened for their ability to inhibit HIV replication using known techniques. For example, HIV virus replication may be monitored using the Cytopathic Effect  
10 (CPE) assay disclosed by Bergeron et al. (J. Virol. 66:5777-5787, 1992). In this assay, the degree of infection is monitored by the appearance of fused cellular membranes ("syncytium"). Alternatively, assays directed to activity of HIV protease may be employed. For example,  
15 the assays and techniques disclosed in the following references may be employed: Ashorn et al., Proc. Natl. Acad. Sci. U.S.A. 87:7472-7476, 1990; Schramm et al., Biochem. Biophys. Res. Commun. 179:847-851, 1991; Sham et al., Biochem. Biophys. Res. Commun. 175:914-919, 1991; and  
20 Roberts et al., Science 248:358-361, 1990. Moreover, the ability of the stable copper(I) complexes of this invention to inhibit HIV replication may be determined by the assay disclosed in Example 5 herein below.

The stable copper(I) complexes of this invention, in  
25 addition to inhibiting HIV replication, may also inhibit replication of other viruses, including human T-cell leukemia (HTLV) I and/or II, human herpes virus, cytomegalo virus (CMV), encephalomyocarditis virus (EMCV), Epstein Barr virus (EBV), human hepatitis virus, Varicella  
30 Zoster virus, Rhinovirus, and rubella virus. One skilled in the art could readily assay the stable copper(I) complexes of this invention for their inhibitory activity with regard to these viruses. For example, Example 15 illustrates the inhibitory affect of stable copper(I)  
35 complexes of this invention on both encephalomyocarditis virus (EMCV) and cytomegalo virus (CMV).

In addition to the biological activity of the stable copper(I) complexes of the present invention, the multi-dentate ligands of this invention also possess biological activity when administered alone as the "free" multi-dentate ligand (i.e., without copper(I)). Such biological activity includes the activities identified above, including anti-viral activity, as well as a preventative agent against gastric tissue damage. Although not intending to be limited to the following theory, when the multi-dentate ligands of this invention are administered as the free ligand, it is believed that they function, at least in part, by scavenging copper(I) to yield the stable copper(I) complex *in vivo*.

The following examples are offered by way of illustration, and not by way of limitation.

#### EXAMPLES

The examples which follow illustrate the preparation and utility of certain exemplary embodiments of the stable copper(I) complexes of the present invention. To summarize the examples that follow: Example 1 illustrates the synthesis of neocuproine copper(I) at a molar ratio of 1:1 and 2:1; Example 2 illustrates the superoxide dismutase (SOD)-mimetic activity of representative copper(I) complexes of this invention (employing a copper(II)-peptide complex as a positive control); Example 3 illustrates the wound healing activity of a representative copper(I) complex of this invention; Example 4 illustrates hair growth activity of a representative copper(I) complex of this invention; Example 5 illustrates inhibition of HIV replication by a representative copper(I) complex of this invention; Example 6 illustrates the activity of a representative "free" multi-dentate ligand of this invention for both wound healing and protection against ethanol-induced gastric mucosal damage; Examples 7 and 8 illustrates the



inhibition of cyclooxygenase-1 and cyclooxygenase-2, respectively, by representative stable copper(I) complexes; Example 9 illustrates the inhibition of 5-lipoxygenase by representative stable copper(I) complexes; Example 10 illustrates the inhibition of leukotriene C<sub>4</sub> synthetase by representative stable copper(I) complexes; Example 11 illustrates the inhibition of elastase by a representative stable copper(I) complex; Example 12 illustrates the inhibition of acetyl coenzyme A synthetase by representative stable copper(I) complexes; Example 13 illustrates the inhibition of HMG-CoA reductase by representative stable copper(I) complexes; Example 14 illustrates the inhibition of HIV-1 activity by various isomers of a representative stable copper(I) complex; Example 15 illustrates the anti-viral activity of representative stable copper(I) complexes and a representative free multi-dentate ligand; Example 16 illustrates inhibition of HIV-1 and HIV-2 proteases by representative stable copper(I) complexes; Example 17 illustrates the inhibition of HIV reverse transcriptase by representative stable copper(I) complexes; and Examples 18 and 19 illustrate the inhibition of Protein Kinase C and various tyrosine kinases, respectively, by representative stable copper(I) complexes.

25

#### Example 1

#### Synthesis of Copper(I)-Neocuproine

Neocuproine hydrate was used as received from Aldrich Chemical Company, having the following properties: mp 161-163°C; <sup>1</sup>H NMR (500MHz, DMSO-d<sub>6</sub>) δ 8.32 (2H, d, J = 8.2), 7.85 (2H, s), 7.60 (2H, d, J = 8.1), 2.79 (6H, s); <sup>13</sup>C NMR (125MHz, DMSO-d<sub>6</sub>) δ 158.0, 144.6, 136.1, 126.4, 125.3, 123.1, 24.9.

A. Neocuproine Copper(I) (1:1)

Cuprous chloride (1.98g, 20.0mmol) was added to a stirred, vacuum-degassed solution of neocuproine hydrate (4.53g, 20.0mmol) in acetonitrile (150mL). This solution  
5 was stirred for 2 hours. The resulting suspension was warmed to boiling and filtered. The filtrate was boiled to a volume of about 100mL. This solution was allowed to cool slowly to give dark red needles: mp280-284°C(decomp.,  
lit. 310-320°C)(Healy et al., J. Chem. Soc. Dalton Trans.  
10 2531, 1985);  $^1\text{H}$  NMR (500MHz, DMSO- $d_6$ )  $\delta$  8.74 (2H, d, J = 8.2), 8.21 (2H, s), 7.95 (2H, d, J = 8.2), 2.38 (6H, s);  $^{13}\text{C}$  NMR (125MHz, DMSO- $d_6$ )  $\delta$  157.6, 142.2, 137.4, 127.1, 125.9, 125.6, 25.1; Anal. calcd. for  $\text{C}_{14}\text{H}_{12}\text{ClCuN}_2$ : C, 54.73; H, 3.94; N, 9.12; Cl, 11.54. Found: C, 54.67; H,  
15 3.89; N, 9.04; Cl, 11.40.

B. Neocuproine Copper(I) (2:1)

A vacuum degassed solution of neocuproine hydrate (4.53g, 20.0mmol) in absolute ethanol (150mL) was added to cuprous chloride (990mg, 10.0mmol) via cannula under an  
20 atmosphere of nitrogen. The resulting bright red solution was stirred at room temperature for 2 hours. This mixture was filtered, to remove a small amount of insoluble matter, and evaporated to give 5.64g (100%) of bright red solid. Recrystallization from aqueous methanol gave very  
25 fine needles: mp231-233°C; UV-vis  $\lambda_{\text{max}}$  (95% ethanol) 207nm ( $\epsilon = 63,750\text{M}^{-1}\text{cm}^{-1}$ ), 226nm ( $\epsilon = 76,250$ ), 272nm ( $\epsilon = 60,000$ ), 454nm ( $\epsilon = 6,750$ ),  $^1\text{H}$  NMR (500MHz, DMSO- $d_6$ )  $\delta$  8.75 (2H, br s), 8.22 (2H, s), 7.96 (2H, br s), 2.40 (6H, s);  $^{13}\text{C}$  NMR (125MHz, DMSO- $d_6$ )  $\delta$  157.6, 142.2, 137.3, 127.1, 125.8,  
30 125.6, 25.0; Anal. calcd. for  $\text{C}_{28}\text{H}_{24}\text{ClCuN}_4$ : C, 65.24; H, 4.69; N, 10.87; Cl, 6.88; Cu, 12.33. Found: C, 65.01; H, 4.73; N, 10.75; Cl, 6.84; Cu, 12.70.

Example 2  
Superoxide Dismutase Mimetic Activity  
of Copper(I) Complex

5       As used herein, compounds which possess activity in a  
superoxide dismutase (SOD) assay are termed "SOD  
mimetics." In this example, representative copper(I)  
complexes of this invention were evaluated for SOD mimetic  
activity as measured by the Xanthine Oxidase/NBT method  
10 (see Oberly and Spitz, Handbook of Methods for Oxygen  
Radical Research, R. Greenwald (ed.), pp. 217-220, 1985;  
Auclair and Voisin, Handbook of Methods for Oxygen Radical  
Research, R. Greenwald (ed.), pp. 123-132, 1985). The  
reactions contained the following: 100 uM Xanthine, 56 uM  
15 NBT (Nitro Blue Tetrazolium), 1 unit of Catalase, 50 mM  
Potassium Phosphate Buffer, pH 7.8. The reaction was  
initiated by the addition of Xanthine Oxidase in  
sufficient quantity to obtain an increase in absorbance at  
560 nm of approximately 0.025/min. in a total volume of  
20 1.7 ml. The Xanthine Oxidase was prepared fresh daily and  
stored on ice until used. All the components of the  
reaction are added except the Xanthine Oxidase and the  
spectrophotometer was adjusted to zero at 560 nm. The  
reaction was initiated by the addition of the Xanthine  
25 Oxidase. All reagents were obtained from Sigma Chemical  
Co.

Measurements of the Absorbance at 560 nm were taken  
at 1-2 minute intervals for at least 16 minutes. The  
control consisted of reactions containing zero copper(I)  
30 complex. The copper(I) complexes tested in this example  
were as follows: bathocuproine disulfonate copper(I)  
("BCDS:Cu(I)"); neocuproine copper(I) ("NC:Cu(I)"); and  
2,2'-biquinoline copper(I) ("BQ:Cu(I)"). As a positive  
control, reactions containing a peptide-copper(II) complex  
35 (i.e., glycyl-L-histidyl-L-lysine:copper(II) or "GHK:Cu"),  
which is a known SOD mimetic (see U.S. Patent No.

4,760,051), were also employed. One unit of SOD activity was taken as that amount of sample in micromoles which inhibits the control reaction with the NBT by 50%. The relative activity is then obtained by comparing the micromoles of copper(I) complex necessary to product a 50% inhibition of the control reactions. The lower the value, the more active the compound is as an SOD mimetic. The results of this experiment are presented in Table 8 below.

10

Table 8SOD-Mimetic Activity of Copper(I) Complexes

<u>Exp. No.</u>	<u>Compound</u>	<u>Copper Ratio</u> <u>(ligand:Cu)</u>	<u>Activity</u> <u>(<math>\mu</math>mol per</u> <u><math>\frac{1}{2}</math>Max. Inhib.)</u>	<u>Relative</u> <u>Activity to</u> <u>Control</u>
1	GHK:Cu(II)	2:1	0.055	---
	BCDS:Cu(I)	2:1	0.034	1.6
2	GHK:Cu(II)	2:1	0.0503	---
	BCDS:Cu(I)	2:1	0.0278	1.8
	BCDS:Cu(I)	2:1	0.0018	28
	NC:Cu(I)	2:1	0.0014	36
3	GHK:Cu(II)	2:1	0.0479	---
	NC:Cu(I)	1:1	0.0018	27
	BQ:Cu(I)	2:1	0.0028	17

15

Example 3Wound Healing Activity  
of Copper(I) Complexes

The subcutaneous implantation of stainless steel wound chambers in rats provides a model for the healing of open cavity wounds. Implantation of these chambers triggers a series of responses which reflect the series of

20

phases involved in wound healing - fibrin clot formation, infiltration of white cells, collagen synthesis, and new blood vessel formation.

This assay involves the implantation of a stainless steel chamber (1 X 2.5 cm cylindrical 312 SS, 20 mesh, with Teflon end caps) on the dorsal mid-line of rats. After one week to allow for encapsulation of the chamber, the chamber on each rat was injected with a 0.2 ml saline solution containing 2.7  $\mu$ mol of the copper(I) complex (i.e., BCDS copper(I) 1:1 or 2:1), or with the same volume of saline (0.2 ml) without the copper(I) complex (i.e., control). Injections were made on days 5, 7, 9, 12, 14, 16 and 19. The chambers were then removed on day 21.

The chambers were lyophilized and the interior contents removed for biochemical analysis. The biochemical parameters examined included the total dry weight, protein content, collagen content (i.e., hydroxyproline content after acid hydrolysis) and glycosaminoglycan content or "GAG" (i.e., uronic acid content after acid hydrolysis).

The protein was determined by the method of Lowry et al. (J. Biol. Chem. 193: 265-275, 1951) using Bovine Serum Albumin (BSA) as a standard. The collagen content was determined by acid hydrolysis and a colorimetric assay for hydroxyproline (Bergman et al., Clin. Chim. Acta 27:347-349, 1970), an amino acid specific for collagen. Glycosaminoglycan content was determined by quantitation of the amount of uronic acid (UA). Aliquots of the homogenate were dissolved in 0.5M NaOH, precipitated and washed with ethanol, and uronic acid was determined by a colorimetric assay using 2-phenylphenol as a reagent (Vilím V., Biomed. Biochem. Acta. 44(11/12s):1717-1720, 1985). Glycosaminoglycan content was expressed as  $\mu$ g of uronic acid per chamber.

The results of this experiment are illustrated in Figure 1. Specifically, BCDS copper(I) at both the 1:1

and 2:1 ratio significantly stimulated the glycosaminoglycan content of the injected chamber. Moreover, BCDS copper(I) at both ratios stimulated the collagen content of the injected chambers. Collagen and glycosaminoglycans are two of the critical extracellular matrix components important for tissue regeneration associated with wound healing.

#### Example 4

##### 10 Stimulation of Hair Growth by Copper(I) Complexes

The following example illustrates the stimulation of hair growth in warm-blooded animals after intradermal injection of a copper(I) complex of this invention.

15 The backs of C3H mice (60 days old, telogen hair growth phase) were closely clipped on day 1 using an electric clipper. A sterile saline solution containing the indicated copper complex was then injected intradermally (i.e., infiltrated under the skin) at two locations within the clipped areas of the mice. Injection at two locations provided two test locations within the clipped area of each mouse. Each injection (0.1 ml) contained the indicated amount of the copper(I) complex (i.e., BCDS copper(I) (1:1) complex at 0.14  $\mu\text{mol}$  and 1.4  $\mu\text{mol}$ ) within a sterile saline solution. A group of saline injected mice (0.1 ml) served as controls. Following injection of the copper(I) complex, indications of hair growth were seen within 10 days. The first visual signs were a darkening of the skin in a circular region surrounding the injection site. The size of this region is generally dose dependent, increasing with an increase in dose. The 0.1 ml injections used in this experiment produced a circle of hair growth measuring approximately 0.5  $\text{cm}^2$  to 5  $\text{cm}^2$  in diameter. Active hair growth occurred between 14-20 days following injection, with a maximum effect seen by day 29. Both the number of mice growing

hair at the injection site and the diameter of the hair growth region were determined at day 21. A positive response was expressed as the number of mice exhibiting hair growth at the injection sites compared to the total number of mice injected in the study. The results of this experiment are presented in Table 9 below.

Table 9  
Hair Growth Activity of BCDS Copper(I) Complex

<u>Amount Injected (<math>\mu\text{mol}</math>)</u>	<u>Growth Area (<math>\text{cm}^2</math>)</u>
0.0 (control)	0.0
0.14	1.35 (Std. Dev. 0.42)
1.4	3.06 (Std. Dev. 0.47)

10

Example 5  
Inhibition of HIV Replication  
of Copper(I) Complex

15 In this experiment, the inhibitory effect of bathocuproine disulfonic acid (BCDS) copper(I) (2:1) complex on phytohemagglutinin (PHA) stimulated peripheral blood mononuclear cells is demonstrated.

20 PHA stimulated peripheral blood mononuclear cells (PBMC) were infected by HIV<sub>IIIB</sub> in the presence of the copper(I) complex identified above and cultured in the presence of the copper(I) complex for two weeks. The extent of HIV replication was assayed at 1 and 2 weeks by a p24 antigen capture ELISA assay. More specifically, 25 PBMC was stimulated with PHA for 24 to 72 hours in basal medium, containing RPMI-1640, 10% fetal bovine serum, and 50  $\mu\text{g/mL}$  gentamicin, and then cultured overnight in the presence of 250 units/mL IL-2. Treated PBMC were pelleted by centrifugation and resuspended to  $0.75 \times 10^6/\text{mL}$  in basal 30 medium with appropriate dilutions of the copper(I) complex or with no copper(I) complex added (i.e., control). To each 0.5 mL aliquot of cells, 0.5 mL of appropriate HIV

dilution was added. The virus-cell mixture was incubated for 2 hours at 37°C in a 5% CO<sub>2</sub> humidified atmosphere. Following the incubation period, the PBMC were washed twice in phosphate-buffered saline. Cells were resuspended in 5 mL to  $7 \times 10^4$  cells/mL in basal medium with (or without) the copper(I) complex. Each cell aliquot was dispensed into four replicate wells of a 48 well tissue culture plate. Cells were fed twice a week with appropriate medium.

At one week and two week culture timepoints the extent of HIV replication was assayed by a p24 antigen capture assay kit (Coulter Corp., Hialeah, Florida). PBMC were treated with buffered detergent to release viral proteins. The cell extract was absorbed to immunoassay titer plates and p24 was detected by binding of a monoclonal anti-p24 antibody coupled to an enzyme. Following the addition of a chromogenic substrate, the amount of p24 was quantified spectrophotometrically.

The results of this experiment are presented in Figure 2. In particular, a 50 uM concentration of the BCDS copper(I) (2:1) complex completely inhibited HIV replication at both week 1 and week 2 at the identified virus dilutions. Furthermore, the 5 uM concentration of BCDS copper(I) (2:1) complex completely inhibited HIV replication at week 2 at the  $10^{-6}$  virus dilution.

#### Example 6

##### Activity of "Free" Multi-Dentate Ligand

This example illustrates the activity of the free multi-dentate ligands of this invention. As used herein, the free ligand is not complexed to the copper(I) ion prior to administration.



A. Inhibition of Ethanol-Induced Gastric Mucosal Damage

Juvenile Sprague-Dawley rats were used in this example. After fasting for 24 hours, the rats were treated by oral gavage with bathocuproine disulfonic acid (BCDS) as the copper(I)-free ligand at various dosages (i.e., 0, 7.6 and 37.6 mg/kg body weight). One hour after BCDS treatment, the animals were challenged with 1 ml of 95% ethanol by oral gavage to cause erosion of the gastric mucosa. As shown in Table 10, BCDS pre-treatment led to a dose-dependent protection against the mucosal damage observed in the control animals.

Table 10  
Effect of BCDS on Ethanol-Induced  
Gastric Mucosal Damage

<u>Dosage</u> <u>mg/kg body weight</u>	<u>Mucosal Damage</u>	
	<u>% of total area</u>	
	<u>Mean</u>	<u>S.E.M.</u>
0.0	45.48	6.94
7.6	32.95	7.49
37.6	23.45	8.18

B. Wound Healing Activity

The BCDS ligand was also examined in the rat wound chamber model as disclosed above in Example 3. The results of this experiment are presented in Table 11.

Table 11  
Effect of BCDS on Wound Healing

<u>mg/injection</u>	<u>ug uronic acid/mg protein</u>
0.0 (control)	28.3 (Std. Dev. $\pm$ 8.7)
1.5	57.6 (Std. Dev. $\pm$ 9.1)
7.5	79.2 (Std. Dev. $\pm$ 10.8)

These results indicate that glycosaminoglycan synthesis is stimulated by administration of the free BCDS ligand.

#### Example 7

##### 5                    Inhibition of Cyclooxygenase-1 by                     Neocuproine and BCDS Copper(I) Complexes (2:1)

Cyclooxygenase is involved in the formation of prostaglandins and thromboxanes by the oxidative  
10 metabolism of arachidonic acid (see Figure 3).

In this experiment, cyclooxygenase-1 from ram seminal vesicles was incubated with arachidonic acid (100  $\mu$ M) for 2 minutes at 37° C in the presence or absence of neocuproine copper(I) (2:1) or BCDS copper(I) (2:1) at  
15 increasing concentrations of neocuproine copper(I) or BCDS copper(I) from 0.3 to 300  $\mu$ M. The assay was terminated by the addition of trichloroacetic acid (TCA), and cyclooxygenase-1 activity was determined by reading the absorbance at 530 nm (Evans et al., "Actions of Cannabis  
20 Constituents on Enzymes of Arachidonate Metabolism: Anti-inflammatory Potential," Biochem. Pharmacol. 36: 2035-2037, 1987; Boopathy and Balasubramanian, "Purification and Characterization of Sheep Platelet cyclooxygenase," Biochem J. 239: 371-377, 1988).

25                    Neocuproine copper(I) (2:1) was found to inhibit cyclooxygenase-1 with an IC<sub>50</sub> of 23  $\mu$ M (see Table 12).. BCDS copper(I) (2:1) complex produced approximately 44% inhibition at a concentration of 300  $\mu$ M. These results demonstrate that the stable copper(I) complexes of this  
30 invention are potent inhibitors of prostaglandin synthesis through inhibition of cyclooxygenase-1.

Table 12  
Inhibition of Cyclooxygenase-1 by  
Stable Copper(I) Complexes

<u>Compound</u>	<u>Conc.</u> ( $\mu$ M)	<u>Percent Inhibition</u> (Mean $\pm$ SEM)		
BCDS Copper(I) (2:1)	300	43.5	$\pm$	1.5
Neocuproine Copper(I) (2:1)	300	77.3	$\pm$	1.5
	30	54.5	$\pm$	0.5
	3.0	15.5	$\pm$	2.5
	0.3	6.5	$\pm$	0.5

5

Example 8  
Inhibition of Cyclooxygenase-2 by  
Neocuproine and BCDS Copper(I) Complexes (2:1)

10       Cyclooxygenase-2, also known as prostaglandin H synthase-2, catalyzes the oxygenation of unesterified precursors to form cyclic endoperoxide derivatives, including prostaglandin H (see Figure 3).

15       In this experiment, cyclooxygenase-2 from sheep placenta, 80 units/tube, was pre-incubated with 1 mM glutathione (GSH), 1 mM hydroquinone, 2.5  $\mu$ M hemoglobin, and either neocuproine copper(I) (2:1) or BCDS copper(I) (2:1) at increasing concentrations of neocuproine copper(I) or BCDS copper(I) from 0.3 to 300  $\mu$ M for 1  
20       minute at 25°C. The reaction was initiated by the addition of arachidonic acid (100  $\mu$ M), and terminated after 20 minutes at 37° C by the addition of TCA. Following centrifugal separation of the precipitated protein, thiobarbiturate was added and cyclooxygenase activity was  
25       determined by absorbance at 530 nm (see Evans et al., *supra*; Boopathy and Balasubramanian, *supra*; O'Sullivan et al., "Lipopolysaccharide Induces Prostaglandin H Synthase-2 in Alveolar Macrophages," *Biochem. Biophys. Res. Commun.* 187:1123-1127, 1992).

Neocuproine copper(I) (2:1) was found to inhibit cyclooxygenase-2 at an estimated IC<sub>50</sub> of 25 $\mu$ M (see Table 13), which is similar to the results of Example 7 with cyclooxygenase-1. BCDS copper(I) (2:1) produced approximately 34% inhibition at the screening concentration of 300  $\mu$ M. These results show that stable copper(I) complexes of this invention are also potent inhibitors of prostaglandin synthesis through inhibition of cyclooxygenase-2.

10

Table 13  
Inhibition of Cyclooxygenase-2 by  
Stable Copper(I) Complexes

<u>Compound</u>	<u>Conc.</u> ( $\mu$ M)	<u>Percent Inhibition</u> (Mean $\pm$ SEM)		
BCDS Copper(I) (2:1)	300	34.0	$\pm$	1.0
Neocuproine Copper(I) (2:1)	300	63.8	$\pm$	0.5
	30	54.0	$\pm$	1.0
	3.0	7.0	$\pm$	1.0
	0.3	6.5	$\pm$	2.5

15

Example 9  
Inhibition of 5-Lipoxygenase by  
Neocuproine and BCDS Copper(I) Complexes (2:1)

20

The 5-lipoxygenase is the principal lipoxygenase in basophils, polymorphonuclear (PMN) leukocytes, macrophages, mast cells, and any organ undergoing an inflammatory response. As illustrated in Figure 3, the action of 5-lipoxygenase leads to the formation of 5-HPETE and 5-HETE, which are precursors to the leukotriene LTB<sub>4</sub> and LTC<sub>4</sub>.

25

In this experiment, 5-lipoxygenase assays were run using a crude enzyme preparation prepared from rat

- basophilic leukemia cells (RBL-1). Neocuproine copper(I) (2:1) or BCDS copper(I) (2:1) at increasing concentrations from 0.3 to 300  $\mu$ M were pre-incubated with the 5-lipoxygenase for 5 minutes at room temperature, and the reaction was initiated by addition of arachidonic acid substrate. After incubation at room temperature for 8 minutes, the reaction was terminated by the addition of citric acid. The levels of 5-HETE were determined by a specific 5-HETE RIA (Shimuzu et al., "Enzyme with Dual Lipoxygenase Activities Catalyzes Leukotriene A<sub>4</sub> Synthesis from Arachidonic Acid," Proc. Natl. Acad. Sci. U.S.A. 81:689-693, 1984; Egan and Gale, "Inhibition of Mammalian 5-Lipoxygenase by Aromatic Disulfides," J. Biol. Chem. 260:11554-11559, 1985).
- Both BCDS copper(I) (2:1) and neocuproine copper (I) (2:1) were found to be inhibitors of 5-lipoxygenase with estimated IC<sub>50</sub>'s of less than 10  $\mu$ M (see Table 14). These results show that stable copper(I) complexes of this invention are potent inhibitors of neutrophil 5-lipoxygenase, thus preventing the accumulation of inflammatory lipid mediators at the sites of inflammation.

Table 14  
Inhibition of 5-Lipoxygenase by  
Stable Copper(I) Complexes

<u>Compound</u>	<u>Conc.</u> ( $\mu$ M)	<u>Percent Inhibition</u> (Mean $\pm$ SEM)
BCDS Copper(I) (2:1)	30	71.3 $\pm$ 2.5
	3.0	29.0 $\pm$ 5.0
	0.3	5.5 $\pm$ 3.5
	0.03	4.0 $\pm$ 1.0
Neocuproine Copper(I) (2:1)	30	99.0 $\pm$ 0.6
	3.0	51.0 $\pm$ 6.0
	0.3	15.5 $\pm$ 2.5
	0.03	7.0 $\pm$ 0.0

Example 10

5                    Inhibition of Leukotriene C<sub>4</sub> Synthetase by  
                  Neocuproine and BCDS Copper(I) Complexes (2:1)

Leukotriene C<sub>4</sub> (LTC<sub>4</sub>) Synthetase is involved in the formation of LTC<sub>4</sub> from LTA<sub>4</sub>, as illustrated in Figure 3, by the addition of a reduced glutathione at the C6 site.

10            In this example, LTC<sub>4</sub> Synthetase was prepared as a crude fraction from rat basophilic leukemia cells (RBL-1). The crude enzyme fraction was incubated with test compounds, LTA<sub>4</sub> methyl ester, albumin (to stabilize the product), and serine borate (to prevent conversion of LTC<sub>4</sub>  
15            to LTD<sub>4</sub>) for 15 minutes at 37° C. The reaction was terminated by the addition of ice cold methanol, and LTC<sub>4</sub> concentration was determined by a specific RIA (Bach et al., "Inhibition by Sulfasalazine of LTC<sub>4</sub> Synthetase and of Rat Liver Glutathione S-Transferases," Biochem.  
20            Pharmacol. 34:2695-2704, 1985; Fitzpatrick et al., "Albumin Stabilizes Leukotriene A<sub>4</sub>," J. Biol. Chem. 257:4680-4683, 1982).

Both BCDS copper(I) (2:1) and neocuproine copper(I) (2:1) were found to be inhibitors of LTC<sub>4</sub> Synthetase with  
25            estimated IC<sub>50</sub>'s of 87 and 285 μM, respectively (see Table 15). These results show that stable copper(I) complexes are potent inhibitors of neutrophil LTC<sub>4</sub> Synthetase, thus preventing the accumulation of inflammatory lipid mediators at the sites of inflammation.

Table 15  
Inhibition of Leukotriene C<sub>4</sub> (LTC<sub>4</sub>) Synthetase by  
Stable Copper(I) Complexes

<u>Compound</u>	<u>Conc.</u> ( $\mu$ M)	<u>Percent Inhibition</u> (Mean $\pm$ SEM)		
BCDS Copper(I) (2:1)	1000	77.8	$\pm$	1.9
	100	51.0	$\pm$	4.0
	10	26.5	$\pm$	1.5
	1	11.0	$\pm$	2.0
Neocuproine Copper(I) (2:1)	1000	71.0	$\pm$	1.9
	100	32.5	$\pm$	0.5
	10	15.0	$\pm$	1.0
	1	9.0	$\pm$	1.0

5

Example 11  
Inhibition of Elastase by  
BCDS Copper(I) (2:1)

10

Proteolysis of various cellular targets by elastase has been implicated in a number of pathologic conditions, including emphysema, rheumatoid arthritis, and psoriasis.

In this experiment, human neutrophil was the source of the elastase. In particular, human neutrophil elastase was prepared in crude form from fresh blood following dextran sedimentation, leukocyte isolation, cell lysis and homogenization of sub-cellular granules containing the elastase. BCDS copper(I) (2:1) was incubated with the enzyme and substrate (methoxysuccinyl-alanyl-alanyl-propyl-valine-4-nitroanalide) for 8 minutes at 25°C. The reaction is terminated by immersing the test tubes in boiling water for 5 minutes. Spectrophotometric analysis of the proteolytic product is measured at 410 nm (Baugh and Travis, "Human Leukocyte Granule Elastase, Rapid

25

Isolation and Characterization," Biochemistry 15:836-841, 1976).

BCDS copper(I) (2:1) was found to inhibit human neutrophil elastase with an estimated IC<sub>50</sub> of 12  $\mu$ M (see Table 16). These results show that stable copper(I) complexes of this invention are potent inhibitors of neutrophil elastase, thus preventing or limiting the breakdown of normal tissue at the sites of inflammation.

10

Table 16

Inhibition of Human Neutrophil Elastase by  
Stable Copper(I) Complexes

<u>Compound</u>	<u>Conc.</u> ( $\mu$ M)	<u>Percent Inhibition</u> (Mean $\pm$ SEM)		
BCDS Copper(I) (2:1)	30	65.8	$\pm$	3.1
	3.0	25.0	$\pm$	5.0
	0.3	18.5	$\pm$	0.5
	0.03	5.5	$\pm$	0.5

15

Example 12

Inhibition of Acetyl Coenzyme A (CoA) Synthetase by  
Neocuproine and BCDS Copper(I) (2:1)

20 In this experiment, the ability of two stable copper(I) complexes, neocuproine copper(I) (2:1) and BCDS copper (I) (2:1), to inhibit certain key enzymes involved in the formation of lipids is demonstrated.

CoA synthetase (yeast) activity was monitored by utilization of a labeled substrate, sodium [3H]acetate (Grayson and WestKaemper, "Stable Analogs of Acyl Adenylates, Inhibition of Acetyl and Acyl (acyl-CoA) CoA Synthetase by Adenosine 5'-alkylphosphates," Life Sci. 43: 437-444, 1988). A reaction buffer including 0.1 M glycine-NaOH (pH 9.0), ATP, and the substrate was pre-

25

30



incubated for 5 minutes at 27°C, followed by addition of 2 nM coenzyme A for an additional 5 minute incubation at 27°C. The reaction was terminated by addition of HCl, and the remaining substrate determined by scintillation counting.

The results of this experiment are presented in Table 17. Both BCDS copper(I) (2:1) and neocuproine copper(I) (2:1) were found to inhibit acetyl CoA synthetase activity.

Table 17  
Inhibition of Acetyl CoA Synthetase by  
Stable Copper(I) Complexes

<u>Compound</u>	<u>IC<sub>50</sub> (μM)</u>
BCDS Copper(I) (2:1)	29
Neocuproine Copper(I) (2:1)	47
Reference compounds:	
Ethyl-5-AMP	60
Lovastatin	>100
Orotic Acid	>100

Both stable copper(I) complexes tested were found to inhibit acetyl CoA synthetase with estimated IC<sub>50</sub>'s of 30-50 μM. These results indicate that the stable copper(I) complexes of this invention may serve as lipid modulating (e.g., lipid lowering) agents.

Example 13  
Inhibition of HMG-CoA Reductase by  
Neocuproine and BCDS Copper(I) (2:1)

In this experiment, HMG-CoA reductase was isolated from rat liver and incubated with [<sup>14</sup>C]HMG-CoA and either neocuproine copper(I) (2:1) or BCDS copper(I) (2:1) for 15

- minutes at 37°C. The reaction is terminated by addition of HCl, and [<sup>14</sup>C] MVA is separated from the intact substrate by column filtration (Kubo and Strott, "Differential Activity of 3-hydroxy-3-methylglutaryl Coenzyme A Reductase in Zones of the Adrenal Cortex," Endocrinology 120: 214-221, 1987; Heller and Gould, "Solubilization and Partial Purification of Hepatic 3-hydroxy-3-methylglutaryl Coenzyme A Reductase," Biochem. Biophys. Res. Comm. 50: 859-865, 1973).
- 10 Testing at 30 µM indicated that both neocuproine copper(I) (2:1) and BCDS copper(I) (2:1) inhibited the HMG-CoA reductase enzyme. The results of this experiment are presented in Table 18.

15 Table 18  
Inhibition of HMG-CoA Reductase by  
Stable Copper(I) Complexes

<u>Compound</u>	<u>IC<sub>50</sub></u>
BCDS Copper(I) (2:1)	>30 µM, <50 µM
Neocuproine Copper(I) (2:1)	>30 µM, <50 µM
Reference compound:	
Lovastatin	>12 nM

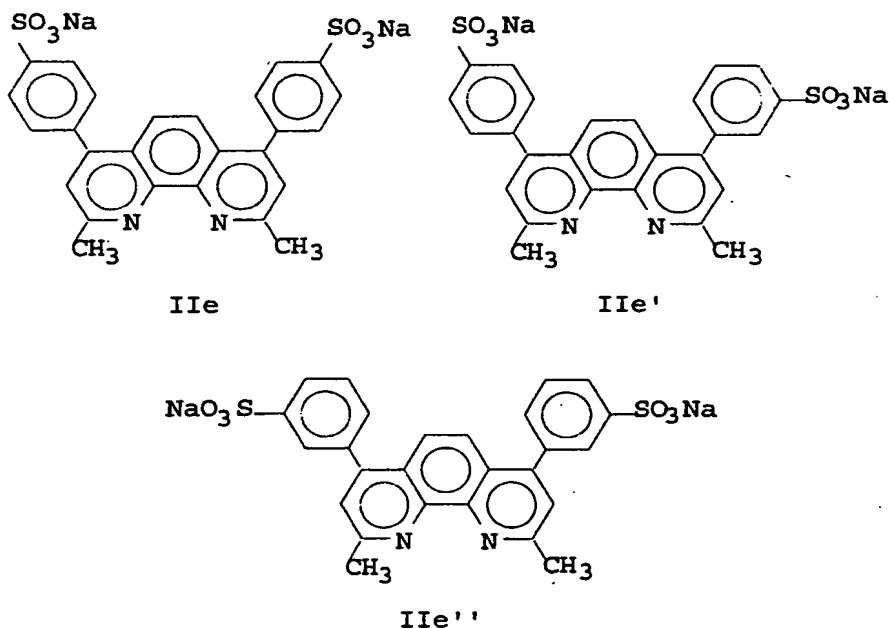
- 20 Both stable copper(I) complexes tested were found to inhibit HMG-CoA reductase with IC<sub>50</sub>'s estimated at greater than 30 µM. These results indicate that the stable copper(I) complexes of this invention may serve as lipid modulating (e.g., lipid lowering) agents.

### Example 14

## Inhibition of HIV-1 Activity by BCDS Copper(I) (2:1) Isomers

5 The experiments presented in this example demonstrate  
the effect on anti-HIV activity of different isomers of  
BCDS copper(I) (2:1). Two experiments utilized p24  
antigen capture as a marker for viral replication, while  
two further experiments utilized reverse transcriptase  
10 activity to monitor the course of infection. The  
infection in all three experiments was performed in  
cultures of human peripheral blood mononuclear cells  
(PBMC) treated with HIV-1.

The positional isomers of BCDS copper(I) employed in  
15 this experiment are identified above as structures IIe,  
IIe' and IIe'', and are set forth below:



Structure IIe is referred to herein as the para-para  
25 ("PP") BCDS isomer since both disulfonic acid/sodium salt

moieties are located in the para position. Similarly, structure IIe' and IIe'' are referred to herein as the meta-para ("MP") and meta-meta ("MM") BCDS isomers, respectively. In addition, a mixture of the PP, MP and MM BCDS isomers was also tested (referred to herein simply as "BCDS"), having a ratio of PP:MP:MM of approximately 5:39:56.

In the first experiment, the anti-HIV activity of BCDS, MP-BCDS and MM-BCDS copper(I) (2:1) at two concentrations (i.e., 10 and 25  $\mu$ M) was compared. These concentrations had been previously determined to be partially and completely effective, respectively, for inhibition of HIV replication by BCDS copper(I).

The same methodology as described above in Example 5 for evaluating inhibition of HIV replication was employed in the experiment. The results of this experiment are present in Table 19.

Table 19

Inhibition of HIV Replication by  
BCDS, MP-BCDS and MM-BCDS Copper(I) (2:1) as Measured by  
p24 Antigen Capture (@1:1000 viral dilution)

	<u>Compound</u>	<u>Week 1</u>		
		<u>p24</u>	<u>(SEM)</u>	<u>% Inhibition</u>
25	Control (infected cells)	30910.00	3770.00	--
	BCDS Copper(I) (10 $\mu$ M)	1959.00	317.16	93.66
	BCDS Copper(I) (25 $\mu$ M)	0.25	0.25	99.99
	MP-BCDS Copper(I) (10 $\mu$ M)	404.50	124.66	98.69
30	MP-BCDS Copper(I) (25 $\mu$ M)	0.50	0.50	99.99
	MM-BCDS Copper(I) (10 $\mu$ M)	346.50	106.27	98.88
	MM-BCDS Copper(I) (25 $\mu$ M)	0.00	0.00	100.00

		<u>Week 2</u>		
<u>Compound</u>		<u>p24</u>	<u>(SEM)</u>	<u>% Inhibition</u>
	Control (infected cells)	10483.80	1109.73	--
	BCDS Copper(I) (10 $\mu$ M)	3286.00	242.36	68.66
5	BCDS Copper(I) (25 $\mu$ M)	0.00	0.00	100.00
	MP-BCDS Copper(I) (10 $\mu$ M)	901.75	277.26	91.40
	MP-BCDS Copper(I) (25 $\mu$ M)	0.00	0.00	100.00
	MM-BCDS Copper(I) (10 $\mu$ M)	549.50	176.25	94.76
	MM-BCDS Copper(I) (25 $\mu$ M)	0.00	0.00	100.00

10

In a second experiment, the activity of BCDS copper(I) and PP-BCDS copper(I) was compared in the manner described above. The results of this experiment are set forth in Table 20. In this experiment the p24 concentrations were lower than in the above experiment. This is due to a different ELISA technique used in this experiment. The standard curve for p24 detection maximizes at 300 pg/ml. Any values over 300 require a kinetic extrapolation to estimate the p24 concentration. Such extrapolation gives a substantial underestimation of the actual p24 concentration. To obtain a more accurate estimate, a series of dilutions of the sample was made to arrive at a reading that is in the middle of the standard curve, and the dilution factor applied to the reading to give the p24 concentrations. This method (which was used in the first experiment, see Table 19 above) while more accurate, yields an overestimate due to the errors of dilution. Nevertheless, the comparisons from one sample to the next in each experiment reflect the inhibitory effects of stable copper(I) complexes tested.

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Table 20  
Inhibition of HIV Replication by  
BCDS and PP-BCDS Copper(I) (2:1) as Measured by  
p24 Antigen Capture (@1:1000 viral dilution)

5				
		<u>Week 1</u>		
	<u>Compound</u>	<u>p24</u>	<u>(SEM)</u>	<u>% Inhibition</u>
	Control (infected cells)	1649.75	29.32	--
	BCDS Copper(I) (10 $\mu$ M)	474.25	41.22	71.25
10	BCDS Copper(I) (25 $\mu$ M)	39.50	6.06	97.61
	PP-BCDS Copper(I) (10 $\mu$ M)	480.00	49.65	70.90
	PP-BCDS Copper(I) (25 $\mu$ M)	34.50	4.57	97.91
15				
		<u>Week 2</u>		
	<u>Compound</u>	<u>p24</u>	<u>(SEM)</u>	<u>% Inhibition</u>
	Control (infected cells)	2256.50	45.93	--
	BCDS Copper(I) (10 $\mu$ M)	1785.00	49.03	20.90
	BCDS Copper(I) (25 $\mu$ M)	22.00	6.38	99.02
	PP-BCDS Copper(I) (10 $\mu$ M)	1915.75	69.75	15.10
20	PM-BCDS Copper(I) (25 $\mu$ M)	33.25	6.60	98.53

In a third experiment, the anti-HIV activity of BCDS, PP-BCDS, MP-BCDS and MM-BCDS copper(I) (2:1) was determined by monitoring the same type of culture (i.e., HIV-1, PBMC) by measuring the reverse transcriptase activity as an infection marker. The PBMC culture conditions for this experiment are described above in Example 5. Following 6 days of incubation, the activity of HIV-1 reverse transcriptase in cellular extracts was determined as a marker for the replication of the virus in culture. The measurement of HIV-1 reverse transcriptase in PBMC cultures may be performed by known techniques (Chattopadhyay et al., "Purification and Characterization of Heterodimeric Human Immunodeficiency Virus Type 1 (HIV-1) Reverse Transcriptase Produced by an In Vitro Processing of p66 with Recombinant HIV-1 Protease," J.

Biol. Chem. 267:14227-14232, 1992). The results of this experiment are presented in Table 21.

Table 21

5

Inhibition of HIV Replication by  
BCDS, PP-BCDS, MP-BCDS and MM-BCDS Copper(I) (2:1)  
as Measured by Reverse Transcriptase Activity

<u>Reverse Transcriptase Activity</u>			
<u>Compound</u>	<u>Conc. (<math>\mu</math>M)</u>	<u>CPM</u>	<u>% Inhibition</u>
None (control)	0	29283	NA
BCDS Copper(I)	0.001	23963	18.17
	0.01	19585	33.12
	0.1	17340	40.78
	1	17623	39.82
	10	4974	83.01
	100	585	98.00
PP-BCDS Copper(I)	0.001	26934	8.02
	0.01	28097	4.05
	0.1	12742	56.49
	1	12247	58.18
	10	1846	93.70
	100	566	98.07
MP-BCDS Copper(I)	0.001	19966	31.82
	0.01	15040	48.64
	0.1	12369	57.76
	1	9880	66.26
	10	1408	95.19
	100	540	98.16
MM-BCDS Copper(I)	0.001	22679	22.55
	0.01	18212	37.81
	0.1	18464	36.95
	1	2085	92.88
	10	583	98.01

In a fourth experiment, inhibition of HIV-1, HIV-2 and SIV, as compared to AZT, was determined for BCDS copper(I), PP-BCDS copper(I), MP-BCDS copper(I) and MM-BCDS copper(I). The experimental conditions described above where employed utilizing Reverse Transcription assay to monitor infection. The results of this experiment are presented in Table 22. It should be noted that the data presented in Table 22 are report in a different format from that of Table 21. In particular, the data of Table 22 represent the calculated EC<sub>50</sub> values. The EC<sub>50</sub> is determined by non-linear regression from inhibition data (such as that presented in Table 21), and extrapolated for the concentration of the test compound required to accomplish a 50% inhibition of reverse transcriptase activity.

Table 22  
Inhibition of HIV-1, HIV-2 and SIV Replication by  
BCDS, PP-BCDS, MP-BCDS and MM-BCDS Copper(I) (2:1)  
as Measured by Reverse Transcriptase Activity

<u>Compound</u>	<u>EC50 (μM)</u>		
	<u>HIV-1</u>	<u>HIV-2</u>	<u>SIV</u>
BCDS Copper(I)	1.7	17.6	4.6
PP-BCDS Copper(I)	0.25	1.2	6.4
MP-BCDS Copper(I)	0.04	12.1	4.3
MM-BCDS Copper(I)	0.13	0.62	6.1
AZT	0.004-0.009	0.0004	0.0066

Example 15  
Anti-Viral Activity of Stable Copper(I) Complexes

This example illustrates that the stable copper (I) compounds of this invention, as well as the free ligands,



have general anti-viral activity. In this experiment, BCDS copper(I) and BCDS alone (i.e., the free ligand) were assayed for the ability to inhibit the murine virus encephalomyocarditis (EMCV) and the cytomegalo virus (CMV).

#### Inhibition of EMCV

Cultures of A<sub>549</sub> cells (human lung) were infected with EMCV for 24-48 hours in the presence of either BCDS copper(I) or BCDS alone. The cells were cultured in DMEM (10% FBS) for 3-4 days prior to use. The medium was then removed, and the cells incubated with sufficient EMCV in serum free DMEM to kill between 30-90% of the cells in the culture. After 2-3 hours of incubation of the cells with EMCV in their presence (or absence) of the test compounds, complete medium (DMEM + 10% FBS) was added and the cells allowed to incubate for 1-2 days in the presence or absence of the test compounds at concentrations ranging from 0.0001-0.0005 M.

The viability of the cultures was then measured by mitochondrial function test (Mossman, J. Immunol. Meth. 65:55-63,1983). The ability of the test compounds to protect the cells from the lethality of the EMCV infection was calculated as a percent protection compared to the mitochondrial activity of parallel, uninfected cells. The results of this experiment are presented in Table 23.

Table 23

#### Inhibition of EMCV by Stable Copper(I) Complex and Free Ligand

30

<u>Conc. (μM)</u>	<u>% Protection</u>	
	<u>BCDS Copper(I) (2:1)</u>	<u>BCDS Ligand</u>
100	30.2	10.7
200	67.2	3.2
400	97.7	22.2
500	153.2	84.2

Inhibition of CMV

Normal Diploid Human Fibroblasts were isolated and cultured with Minimal Essential Medium (MEM) containing Earles balanced salts and supplemented with 10% Fetal Bovine Serum (FBS). Cytomegalo virus (CMV) was added to the cultures in the presence or absence of BCDS and BCDS copper(I) (2:1). Five cultures were employed in each test group, with the exception of the uninfected cell groups which utilized 8 cultures. The uninfected cell groups were used to ensure that antiviral activity was achieved in the absence of any direct cytotoxic effect of the test compounds.

After one week of incubation, cellular viability (i.e., mitochondrial function) was determined, and the ability of the test compounds to prevent the cytopathic effect (CPE) of the virus was calculated as percent protection by the following formula:

$$\% \text{ Protection} = (V_t - V_v) / (V_u - V_v) \times 100$$

where  $V_t$  represents viability of the test culture,  $V_v$  represents the viability of culture with virus alone, and  $V_u$  represents the viability of uninfected cells.

The results of this experiment are presented in Table 24. No cytotoxic effects were observed on the uninfected compounds treated with the test compounds.

Table 24

Inhibition of CMV by  
Stable Copper(I) Complex and Free Ligand

30

<u>% Protection (SEM)</u>		
<u>Conc. (<math>\mu</math>M)</u>	<u>BCDS Copper(I) (2:1)</u>	<u>BCDS Ligand</u>
25	13.1 (7.7)	34.2 (8.1)
100	117.3 (13.3)	35.4 (6.2)
250	92.9 (5.2)	23.6 (8.3)

Example 16Inhibition of HIV-1 and HIV-2 Proteases by  
Stable Copper(I) Complexes

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This example illustrates the ability of stable copper(I) complexes of this invention to inhibit HIV-1 and HIV-2 proteases.

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HIV-1 Protease <sup>125</sup>I-SPA Assay

In this experiment, SPA beads (Scintillation Proximity Assay) were coupled with a peptide substrate to assay for HIV-1 protease. The substrate was a 12 residue peptide with the following sequence:

15

AcN-Tyr-Arg-Ala-Arg-Val-Phe-Phe-Val-Arg-Ala-Ala-Lys-COOH

The peptide was monoiodinated on the terminal tyrosine residue, biotinylated through the ε-amino group on the terminal lysine, and linked to the SPA bead via a streptavidin link.

20

HIV-1 protease cleaves the peptide substrate at the Phe-Phe bond, releasing the <sup>125</sup>I-fragment from the bead. Once the peptide is cleaved, it can no longer stimulate the scintillant in the SPA bead and the signal is reduced. The rate of reduction is proportional to the activity of the HIV-1 protease. Recombinant HIV-1 protease, affinity purified for kinetic and assay studies, was used in this experiment.

25

30

Two types of controls were conducted with this assay, one without enzyme to test for possible scintillation quenching by the test compound (i.e., BCDS copper(I) (2:1)), and another positive control with acetyl pepstatin. At concentrations 10 times that used in the

35

assay, there was no quench detected in the presence of BCDS copper(I) (2:1).

The results of this experiment are presented in Table 25. The data presented is the mean  $\pm$  SD of the percent inhibition relative to a no enzyme control reaction. As discussed above, the IC<sub>50</sub> was estimated from the point at which the dose inhibition line crossed the 50% inhibition line. The estimated IC<sub>50</sub> with this HIV-1 protease assay was 11 $\mu$ M.

10

Table 25  
Inhibition of HIV-1 Protease by  
Stable Copper(I) Complexes

<u>Compound</u>	<u>Conc.</u> ( $\mu$ M)	<u>Percent Inhibition</u> (Mean $\pm$ SEM)		
BCDS Copper(I) (2:1)	25	86.7	$\pm$	2.1
	10	45.2	$\pm$	2.3
	5	17.6	$\pm$	2.3
	2	12.2	$\pm$	1.8
	1	8.5	$\pm$	4.6
	0.5	1.8	$\pm$	0.6
	0.1	0.0	$\pm$	1.9
Reference Compound:				
Acetyl Pepstatin	0.5	67.4	$\pm$	1.1
	0.25	50.1	$\pm$	0.4
	0.1	28.4	$\pm$	7.7
	0.05	16.6	$\pm$	0.5
	0.025	10.2	$\pm$	2.6
	0.01	2.4	$\pm$	3.5

15

#### HIV-2 Protease <sup>125</sup>I-SPA Assay

As in the above experiment, SPA beads were coupled with a peptide substrate to assay for HIV-2 protease. The substrate was the 12 residue peptide identified above and

20

moniodinated on the terminal tyrosine residue, biotinylated through the  $\epsilon$ -amino group on the terminal lysine, and linked to the SPA bead via a streptavidin link.

5 HIV-2 protease cleaves the peptide substrate at the Phe-Phe bond, releasing the  $^{125}\text{I}$ -fragment from the bead. Once the peptide is cleaved, it can no longer stimulate the scintillant in the SPA bead and the signal is reduced. The rate of reduction is proportional to the activity of  
10 the HIV-2 protease. Recombinant HIV-2 protease, affinity purified for kinetic and assay studies, was used in this experiment. HIV-2 protease has about 50% sequence homology with HIV-1 protease, and is similar to simian immunodeficiency virus (SIV) protease.

15 Two types of control assays were again run, one without enzyme and the other using acetyl pepstatin as a positive control.

The results of this experiment are presented in Table 26. The data presented is the mean  $\pm$  SD of the percent  
20 inhibition relative to a no enzyme control reaction. The  $\text{IC}_{50}$  was estimated from the point at which the dose inhibition line crossed the 50% inhibition line. The estimated  $\text{IC}_{50}$  with this HIV-2 protease assay was  $10\mu\text{M}$ .

Table 26  
Inhibition of HIV-2 Protease by  
Stable Copper(I) Complexes

<u>Compound</u>	<u>Conc.</u> ( $\mu$ M)	<u>Percent Inhibition</u> (Mean $\pm$ SEM)		
BCDS Copper(I) (2:1)	25	51.9	$\pm$	5.5
	10	49.6	$\pm$	2.9
	5	32.2	$\pm$	2.8
	2	14.1	$\pm$	1.1
	1	5.3	$\pm$	0.9
	0.5	0.8	$\pm$	5.3
	0.1	2.5	$\pm$	0.6
Reference Compound:				
Acetyl Pepstatin	5.0	88.8	$\pm$	0.5
	2.5	70.6	$\pm$	2.3
	1.0	45.5	$\pm$	1.8
	0.5	37.2	$\pm$	0.3
	0.25	19.9	$\pm$	5.6
	0.1	4.3	$\pm$	12.3

5

Example 17  
Inhibition of HIV Reverse Transcriptase by  
Stable Copper(I) Complexes

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This example illustrates the ability of a stable copper(I) complex of this invention, BCDS copper(I) (2:1), to inhibit HIV reverse transcriptase activity.

As in Example 16 above, SPA (Scintillation Proximity Assay) beads were used to assay for the reverse transcriptase activity. The reverse transcriptase (10  $\mu$ L) was incubated with the 3H-deoxyribonucleotides (10  $\mu$ L), the DNA primer linked to biotin (10  $\mu$ L), and the RNA template. After incubation at 37° for 20 minutes, the reaction was stopped and the labeled product was recovered

20

by addition of the SPA beads coupled to streptavidin which binds to the biotin linked DNA primer.

The extent of the reaction was determined by scintillation counting. Increasing concentrations of BCDS copper(I) (2:1) were added and the extent of the reaction determined by the method described above.

The results of this experiment are presented in Table 27. The data show the mean  $\pm$  SD of the percent inhibition relative to a no test compound control reaction. The IC<sub>50</sub> is estimated from the point at which the dose inhibition like crosses the 50% inhibition line. The estimated IC<sub>50</sub> was 11 $\mu$ M.

Table 27  
Inhibition of HIV Reverse Transcriptase by  
Stable Copper(I) Complexes

Compound	Conc. ( $\mu$ M)	Percent Inhibition (Mean)
BCDS Copper(I) (2:1)	25	63.2
	10	35.4
	5	26.8
	2	27.0
	1	9.8
	0.5	0.2
	0.1	1.8

Example 18  
Inhibition of Protein Kinase C by  
Stable Copper(I) Complexes

This example illustrates the ability of the representative stable copper(I) complexes, BCDS copper(I) (2:1) and neocuproine copper(I) (2:1), to inhibit enzymes involved in intracellular signal transduction. The

enzymes tested in this experiment were various protein kinase C isozymes [and protein tyrosine kinases specific for growth factors and cytokines].

#### 5                    Protein Kinase C (non-selective) Assay

In this experiment, the reaction mixture included 20 mM Tris-HCl, pH 7.4, [32P]-ATP, phosphatidylserine, partially purified PKC from rat brain, and one of the test compounds (Hunnun, et al. "Activation of Protein Kinase C by Triton X-100 Mixed Micelles Containing Diacylglycerol and Phosphatidylserine," J. Biol. Chem. 260:10039-10043, 1985; Jeng, et al., "Purification of Stable Protein Kinase C from Mouse Brain Cytosol by Specific Ligand Elution Using Fast Protein Liquid Chromatography," Cancer. Res. 46:1966-1971, 1986). Following a 10 minute incubation, 25 ul aliquots are removed, spotted on phosphocellulose paper, washed three times in cold phosphoric acid, dried, and counted to determine phosphorylated product. The results of this experiment are presented in Table 28.

20

Table 28  
Inhibition of Protein Kinase C (non-selective) by  
Stable Copper(I) Complexes

<u>Compound</u>	<u>Conc.</u> ( $\mu$ M)	<u>Percent Inhibition</u> (Mean $\pm$ SEM)		
BCDS Copper(I) (2:1)	300	87.5	$\pm$	2.7
	30	9.5	$\pm$	4.5
	3.0	7.5	$\pm$	4.5
	0.3	2.5	$\pm$	4.5
Neocuproine Copper(I) (2:1)	300	62.0	$\pm$	1.9
	30	22.0	$\pm$	7.0
	3.0	6.0	$\pm$	4.0
	0.3	-12.0	$\pm$	2.0



### Protein Kinase C $\alpha$ Assay

Protein Kinase C $\alpha$  is one of the major protein kinase C isoforms. Protein kinase C is a family of serine/threonine protein kinases that mediate the actions of a wide variety of growth factor, hormone, and neurotransmitter action.

In this experiment, protein Kinase C $\alpha$  was purified to homogeneity from rat brain using a modification of a the published procedure(3). The purity of the isolated PKC $\alpha$  was confirmed by SDS/polyacrylamide gel electrophoresis and isoform-specific antibodies. The enzyme was pre-incubated with the test compounds, and its activity is measured by the ability of the enzyme to phosphorylate histone H1 in the absence and presence of calcium, phosphatidylserine, diolein and [32P]ATP. Following a 5 minute incubation, the reaction was terminated by the addition of acetic acid, 50 ul aliquots are removed, spotted on phosphocellulose paper, washed three times in water, dried, and counted to determine phosphorylated product. The data presented in Table 29 show that the addition of the stable copper(I) complexes inhibit the activity of Protein Kinase C $\alpha$ .

Table 29  
Inhibition of Protein Kinase C $\alpha$  by  
Stable copper(I) Complexes

<u>Compound</u>	<u>Conc.</u> ( $\mu$ M)	<u>Percent Inhibition</u> (Mean $\pm$ SEM)
BCDS Copper(I) (2:1)	100	88.3 $\pm$ 0.6
	10	18.0 $\pm$ 2.0
	1.0	0.0 $\pm$ 3.0
	0.1	-4.5 $\pm$ 6.5

Neocuproine Copper(I) (2:1)	100	87.5	±	1.8
	10	23.6	±	4.5
	1.0	-1.5	±	3.5
	0.1	-5.0	±	1.0

#### Protein Kinase C $\beta$ Assay

Protein Kinase C $\beta$  is another major protein kinase C isoforms. Protein kinase C is a family of  
5 serine/threonine protein kinases that mediate the actions of a wide variety of growth factor, hormone, and neurotransmitter action.

In this experiment, Protein Kinase C $\beta$  (which includes  $\beta$ I and  $\beta$ II forms) was purified to homogeneity from rat  
10 brain using a modification of a published protocol (Woodgett and Hunter, "Isolation and Characterization of Two Distinct Forms of Protein Kinase C," J. Biol. Chem. 262:4836-4848, 1987). The purity of the isolated PKC $\alpha$  was confirmed by SDS/polyacrylamide gel electrophoresis and  
15 isoform-specific antibodies. The enzyme was pre-incubated with test compounds, and its activity is measured by the ability of the enzyme to phosphorylate histone H1 in the absence and presence of calcium, phosphatidylserine, diolein and [32P]ATP. Following a 5 minute incubation,  
20 the reaction was terminated by the addition of acetic acid, 50  $\mu$ l aliquots are removed, spotted on phosphocellulose paper, washed three times in water, dried, and counted to determine phosphorylated product.

The data presented in Table 30 show that the addition  
25 of the stable copper(I) complexes inhibit the activity of Protein Kinase C $\beta$ .

Table 30  
Inhibition of Protein Kinase C $\beta$  by  
Stable Copper(I) Complexes

<u>Compound</u>	<u>Conc.</u> ( $\mu$ M)	<u>Percent Inhibition</u>	
		(Mean $\pm$ SEM)	
BCDS Copper(I) (2:1)	100	96.8	$\pm$ 2.0
	10	20.0	$\pm$ 2.0
	1.0	3.5	$\pm$ 6.5
	0.1	6.5	$\pm$ 4.5
Neocuproine Copper(I) (2:1)	100	84.5	$\pm$ 1.9
	10	25.5	$\pm$ 1.5
	1.0	4.0	$\pm$ 7.0
	0.1	3.5	$\pm$ 6.5

5

#### Protein Kinase Cy Assay

Protein Kinase Cy is another major protein kinase C isoform. Protein kinase C is a family of serine/threonine protein kinases that mediate the actions of a wide variety of growth factor, hormone, and neurotransmitter action.

In this experiment, Protein Kinase Cy was purified from insect cells expressing a baculovirus recombinant rabbit brain protein kinase Cy isoform. The enzyme was pre-incubated with the test compounds, and its activity was measured by the ability of the enzyme to phosphorylate histone H1 in the absence and presence of calcium, phosphatidylserine, diolein and [32P]ATP. Following a 5 minute incubation, the reaction was terminated by the addition of acetic acid, 50  $\mu$ l aliquots were removed, spotted on phosphocellulose paper, washed three times in water, dried, and counted to determine phosphorylated product.

The data presented in Table 31 show that the addition of the stable copper(I) complexes inhibit the activity of Protein Kinase Cy.

Table 31  
Inhibition of Protein Kinase Cy by  
Stable Copper(I) Complexes

<u>Compound</u>	<u>Conc. (<math>\mu</math>M)</u>	<u>Percent Inhibition</u>
BCDS Copper(I) (2:1)	100	99
	10	51
	1.0	21
	0.1	5
Neocuproine Copper(I) (2:1)	100	97
	10	40
	1.0	28
	0.1	17

5

The data in Tables 28-31 was used to determine the 50% inhibitory dose ( $IC_{50}$ ) of each enzyme. This data is presented below in Table 32. These results show that the stable copper(I) complexes of this invention are potent inhibitors of Protein Kinase C.

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Table 32  
Inhibition of Protein Kinase C Isoforms by  
Stable Copper(I) Complexes

<u>Protein Kinase C</u>	<u><math>IC_{50}</math> (<math>\mu</math>M)</u>	
<u>Isoform</u>	<u>BCDS</u>	<u>Neocuproine</u>
	<u>Copper(I) (2:1)</u>	<u>Copper(I) (2:1)</u>
Non-selective	97	145
Ca	28	25
c $\beta$	25	25
Cy	8.8	15

Example 19Inhibition of Protein Tyrosine Kinases by  
Stable Copper(I) Complexes

5        This example illustrates the ability of the  
representative stable copper(I) complexes, BCDS copper(I)  
(2:1) and neocuproine copper(I) (2:1), to inhibit enzymes  
involved in intracellular signal transduction. The  
enzymes tested in this experiment were protein tyrosine  
10    kinases specific for growth factors and cytokines.

Epidermal Growth Factor (EGF)Receptor Tyrosine Kinase (human recombinant) Assay

The binding of EGF or TGF- $\alpha$  (Transforming Growth  
15    Factor  $\alpha$ ) to the EGF receptor results in activation of the  
tyrosine kinase portion of the receptor. This kinase  
phosphorylates several cytosolic proteins which lead to  
induction of intracellular signaling pathways eventually  
leading to cell mitogenesis and in some cases cellular  
20    transformation. Inhibition of the EGF tyrosine kinase is  
useful for chemotherapy for malignant cells.

In this experiment, a recombinant form of the human  
Epidermal Growth Factor Tyrosine Kinase domain was assayed  
(Geissler et al., "Thiazolidine-Diones: Biochemical and  
25    Biological Activity of a Novel Class of Tyrosine Protein  
Kinase Inhibitors," J. Biol. Chem. 165:22255-22261, 1990;  
Wedegartner and Gill, "Activation of the Purified Protein  
Kinase Domain of the Epidermal Growth Factor Receptor," J.  
Biol. Chem. 264:11346-11353, 1989; Yaish et al., "Blocking  
30    of EGF-dependent Cell Proliferation by EGF-Receptor Kinase  
Inhibitors," Science 242:933-935, 1988).

The kinase assay measures the activity of the 69kD  
kinase domain by employing an immobilized synthetic  
polypeptide as a substrate. Following a 10 minute  
35    reaction, phosphorylated tyrosine residues were detected  
by incubation with a monoclonal anti-phosphotyrosine

antibody. Bound anti-phosphotyrosine antibody was quantitated by incubation with a biotin-linked anti-mouse IgG, followed by streptavidin linked  $\beta$ -galactosidase enzyme. Fluorescence resulting from conversion of  
 5 fluoroscein-di- $\beta$ -galactoside to fluorescein was measured. The results of this experiment are presented in Table 33.

Table 33  
Inhibition of Epidermal Growth Factor (EGF) Receptor  
Tyrosine Kinase (human recombinant) by  
Stable Copper(I) Complexes

10

<u>Compound</u>	<u>Conc.</u> ( $\mu$ M)	<u>Percent Inhibition</u> (Mean $\pm$ SEM)		
BCDS Copper(I) (2:1)	10	102.3	$\pm$	2.3
	1	40.0	$\pm$	3.6
	0.1	11.7	$\pm$	2.7
	0.01	0.3	$\pm$	2.4
Neocuproine Copper(I) (2:1)	10	96.7	$\pm$	1.0
	1	43.7	$\pm$	5.5
	0.1	12.0	$\pm$	4.0
	0.01	-5.7	$\pm$	0.9

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p56<sup>lck</sup> Tyrosine Kinase Assay

The lck tyrosine kinase is a member of the src family of cytoplasmic tyrosine kinases. It is expressed only in T-lymphocytes and NK cells. The p56<sup>lck</sup> Tyrosine Kinase is a 56 kD protein that is found associated with the  
 20 cytoplasmic side of the plasma membrane of these cells. It is responsible of transmission of the IL-2 signal leading to T-lymphocyte activation. The binding of IL-2 to specific IL-2 receptors leads to activation of the p56 tyrosine kinase. In addition, the p56<sup>lck</sup> Tyrosine Kinase  
 25 has been found to function in signal transduction for antigen activated CD4 and CD8 T-cell receptors.

In this experiment, the p56<sup>lck</sup> Tyrosine Kinase was purified from bovine thymus. The kinase assay measures the activity of the 69kD kinase domain by employing an immobilized synthetic polypeptide as a substrate. The

5 test compounds were pre-incubated with the enzyme of 15 minutes. Following a 10 minute reaction with 100 uM ATP, phosphorylated tyrosine residues are detected by incubation with a monoclonal anti-phosphotyrosine antibody. Bound anti-phosphotyrosine antibody was

10 quantitated by incubation with a biotin-linked anti-mouse IgG, followed by streptavidin linked  $\beta$ -galactosidase enzyme. Fluorescence resulting from conversion of fluoroscein-di- $\beta$ -galactoside to fluorescein was measured (Hatekeyama et al., "Interaction of the IL-2 Receptor with

15 the src-Family Kinase p56<sup>lck</sup>: Identification of Novel Intermolecular Association," Science 252:1523-1528, 1991; Caron et al., "Structural Requirements for Enhancement of T-cell Responsiveness by the Lymphocyte Specific Tyrosine Protein Kinase p56<sup>lck</sup>," Mol. Cell Biol. 12:2720-2729,

20 1992; Cheng et al., "A Synthetic Peptide Derived from p34cdc2 is a Specific and Efficient Substrate of src-Family Tyrosine Kinases." J. Biol. Chem. 267:9248-9256, 1992).

Both the BCDS copper(I) and neocuproine copper(I)

25 complexes were found to be potent inhibitors of the kinase activity. The results of this experiment are presented in Table 34.

Table 34

30 Inhibition of p56<sup>lck</sup> Tyrosine Kinase Activity  
by Stable Copper(I) Complexes

Compound	Conc.	Percent Inhibition		
	( $\mu$ M)	(Mean $\pm$ SEM)		
BCDS Copper(I) (2:1)	10	97.5	$\pm$	1.7
	1	19.5	$\pm$	1.5

	0.1	-3.5	±	0.5
	0.01	2.5	±	7.5
Neocuproine Copper(I) (2:1)	10	90.0	±	2.3
	1	19.5	±	0.5
	0.1	-8.0	±	2.0
	0.01	-1.0	±	6.0

#### p59<sup>fyn</sup> Tyrosine Kinase Assay

The fyn tyrosine kinase is also a member of the src family of non-receptor linked cytoplasmic tyrosine kinases. The p59<sup>fyn</sup> Tyrosine Kinase is responsible for mediating signal transduction through the T-cell receptor (TCR). This receptor is responsible for a signal cascade leading to lymphokine secretion and cell proliferation. The p59<sup>fyn</sup> Tyrosine Kinase is also one of several kinases associated with the B-cell receptor.

In this experiment, the p59<sup>fyn</sup> Tyrosine Kinase was purified from bovine thymus. The kinase assay measures the activity of the 69kD kinase domain by employing an immobilized synthetic polypeptide as a substrate. The test compounds are preincubated with the enzyme of 15 minutes. Following a 10 minute reaction with 100 uM ATP, phosphorylated tyrosine residues are detected by incubation with a monoclonal anti-phosphotyrosine antibody. Bound anti-phosphotyrosine antibody is quantitated by incubation with a biotin-linked anti-mouse IgG, followed by streptavidin linked  $\beta$ -galactosidase enzyme. Fluorescence resulting from conversion of fluoroscein-di- $\beta$ -galactoside to fluorescein is measured (Cooke et al., "Regulation of T-cell Receptor Signaling by a src Family Protein Tyrosine Kinase p59<sup>fyn</sup>," Cell 65:281-291, 1991; Grassman et al, "Protein Tyrosine Kinase p59<sup>fyn</sup> is Associated with the T-cell Receptor CD3 Complex in Functional Human Lymphocytes," Eur. J. Immunol. 22:283-286, 1992; Appleby et al., "Defective T-cell Receptor



Signaling in Mice Lacking the Thymic Isoform of p59<sup>fyn</sup>, "Cell 70:751-763, 1992). Both the BCDS copper(I) and neocuproine copper(I) complexes were found to be potent inhibitors of the kinase activity. The results of this experiment are presented in Table 35.

Table 35  
Inhibition of p59<sup>fyn</sup> Tyrosine Kinase Activity by  
Stable Copper(I) Complexes

10

<u>Compound</u>	<u>Conc.</u> ( $\mu$ M)	<u>Percent Inhibition</u> (Mean $\pm$ SEM)		
BCDS Copper(I) (2:1)	10	99.0	$\pm$	2.7
	1	38.0	$\pm$	5.0
	0.1	20.5	$\pm$	0.5
	0.01	2.0	$\pm$	8.0
Neocuproine Copper(I) (2:1)	10	91.0	$\pm$	3.0
	1	25.5	$\pm$	1.5
	0.1	1.0	$\pm$	6.0
	0.01	2.5	$\pm$	4.5

The data in Tables 33-35 were used to determine the 50% inhibitory dose (IC<sub>50</sub>) of each protein tyrosine kinase tested. This data is shown in Table 36. These results show that the stable copper(I) complexes of this invention are potent inhibitors of this class of tyrosine kinase.

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Table 36  
Inhibition of Protein Tyrosine Kinases by  
Stable Copper(I) Complexes

<u>Protein Tyrosine Kinase</u>	<u>IC<sub>50</sub> (μM)</u>	
	<u>BCDS</u>	<u>Neocuproine</u>
	<u>Copper(I) (2:1)</u>	<u>Copper(I) (2:1)</u>
EGF Receptor Tyrosine Kinase	1.3	1.4
p56 <sup>lck</sup> Tyrosine Kinase	2.4	2.7
p59 <sup>fyn</sup> Tyrosine Kinase	1.5	2.4

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From the foregoing, it will be appreciated that, although specific embodiments of the invention have been described herein for purposes of illustration, various  
10 modifications may be made without deviating from the spirit and scope of the invention. Accordingly, the invention is not to be limited except as by the appended claims.

Claims

1. Use of a stable copper(I) complex as an active therapeutic substance.

2. A composition comprising a stable copper(I) complex in combination with a pharmaceutically acceptable carrier or diluent.

3. The composition of claim 2 wherein the stable copper(I) complex is (2,9-dimethyl-4,7-diphenyl-1,10-phenanthroline disulfonic acid disodium salt) copper(I) (2:1).

4. The composition of claim 3, wherein the stable copper(I) complex is a single isomer of (2,9-dimethyl-4,7-diphenyl-1,10-phenanthroline disulfonic acid disodium salt) copper(I) (2:1).

5. The composition of claim 2 wherein the stable copper(I) complex is (2,9-dimethyl-1,10-phenyl-1,10-phenanthroline) copper(I) (2:1).

6. A method for enhancing wound healing in a warm-blooded animal, comprising administering to the animal an effective amount of a stable copper(I) complex.

7. A method for enhancing or restoring the resistance of a warm-blooded animal to oxidative or inflammatory damage associated with reactive oxygen species, comprising administering to the animal an effective amount of a stable copper(I) complex.

8. A method for treating inflammation in a warm-blooded animal, comprising administering to the animal an effective amount of a stable copper(I) complex.

9. A method for modulating lipid metabolism in a warm-blooded animal, comprising administering to the animal an effective amount of a stable copper(I) complex.

10. A method for stimulating the growth of hair in a warm-blooded animal, comprising administering to the animal an effective amount of a stable copper(I) complex.

11. A method for modulating signal transduction in a warm-blooded animal, comprising administering to the animal an effective amount of a stable copper(I) complex.

12. A method for inhibiting Protein Kinase C in a warm-blooded animal, comprising administering to the animal an effective amount of a stable copper(I) complex.

13. A method for inhibiting a protein tyrosine kinase in a warm-blooded animal, comprising administering to the animal an effective amount of a stable copper(I) complex.

14. A method for inhibiting viral replication in a warm-blooded animal, comprising administering to the animal an effective amount of a stable copper(I) complex.

15. The method of claim 14, wherein the virus is selected from the group consisting of human T-cell leukemia I and/or II, human herpes virus, cytomegalo virus, encephalomyocarditis virus, Epstein Barr virus, human hepatitis virus, Varicella Zoster virus, Rhinovirus, and rubella virus.

16. The method of claim 14, wherein the virus is human immunodeficiency virus.

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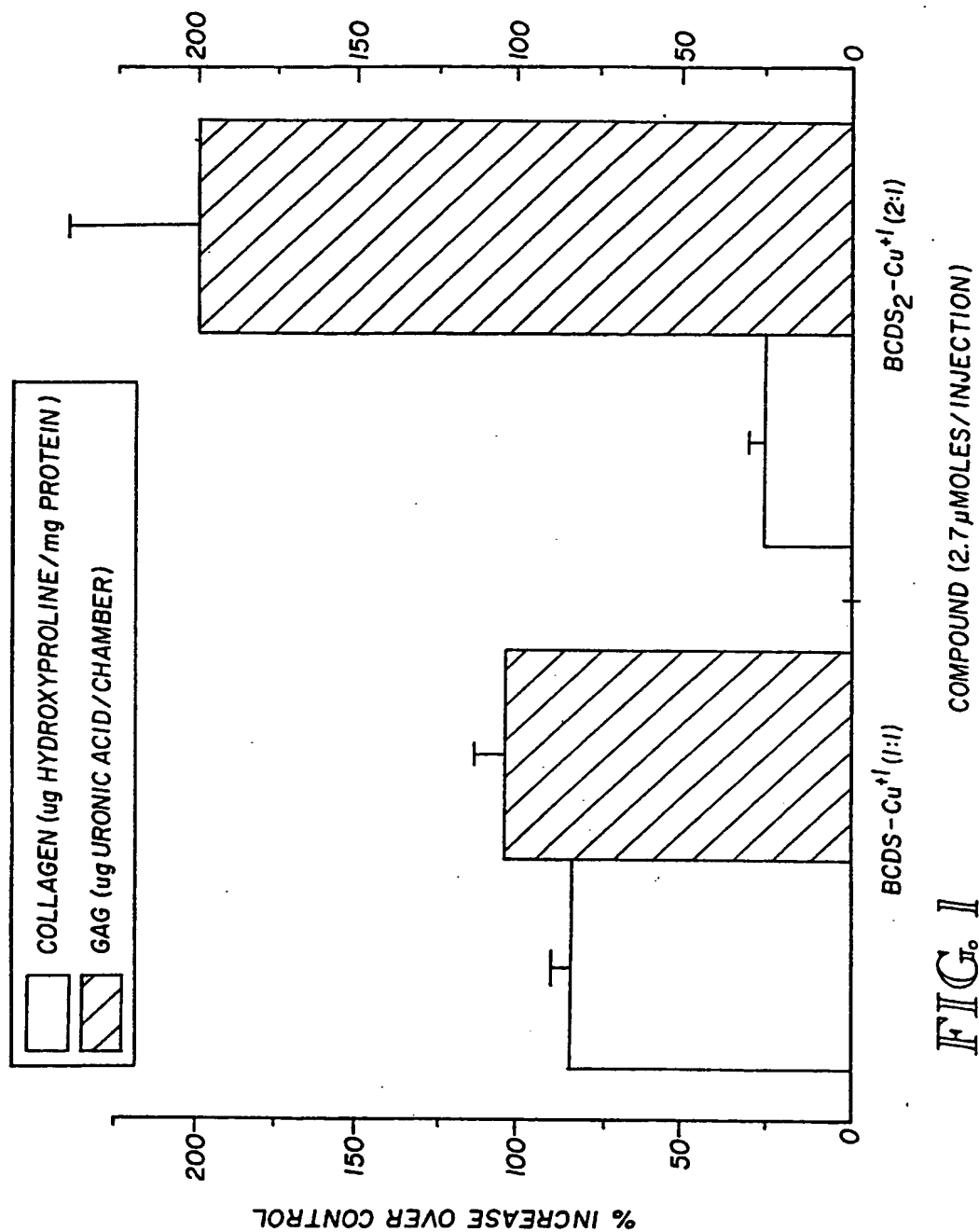


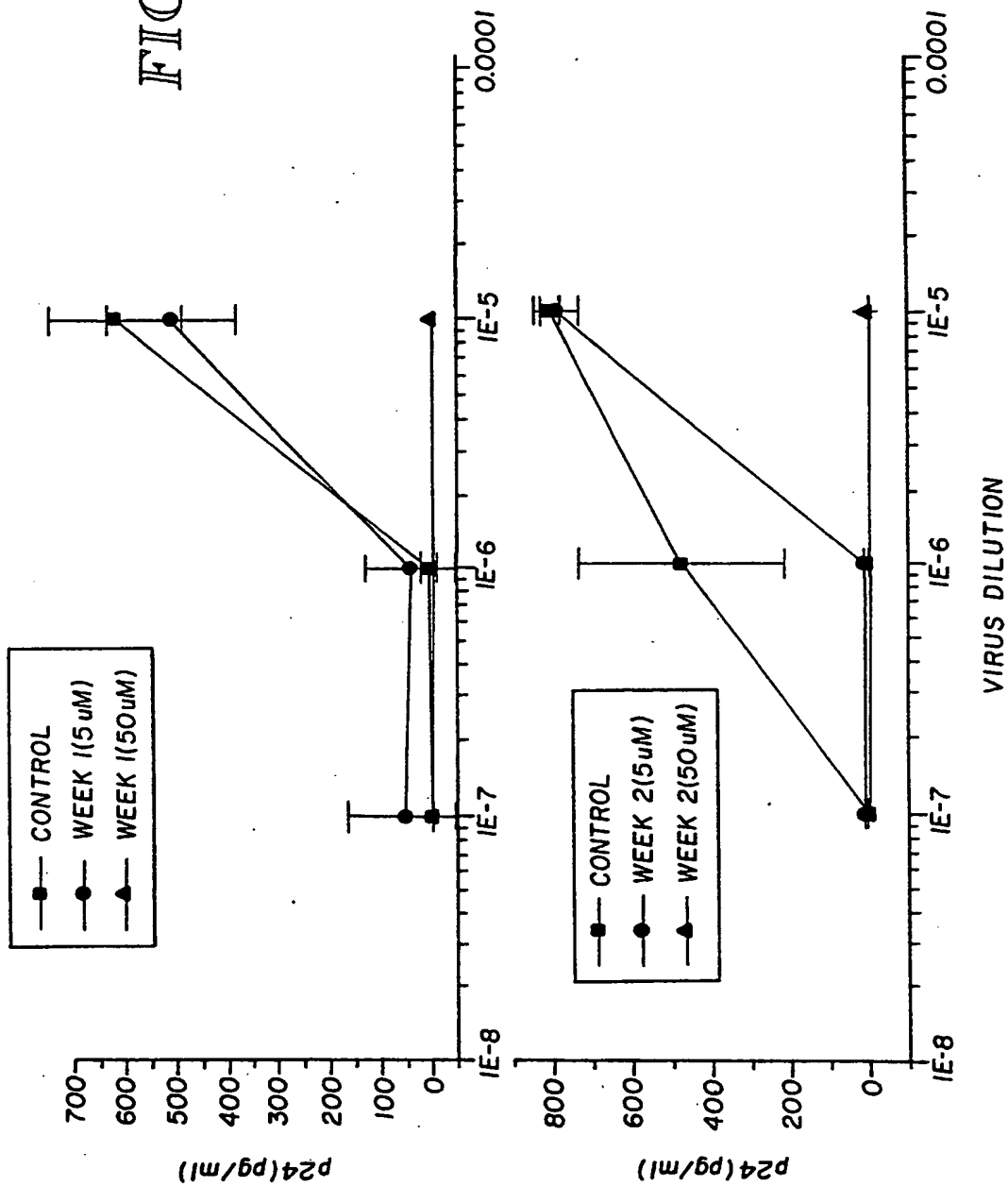
FIG. 1

COMPOUND (2.7 μMOLES/INJECTION)

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FIG. 2



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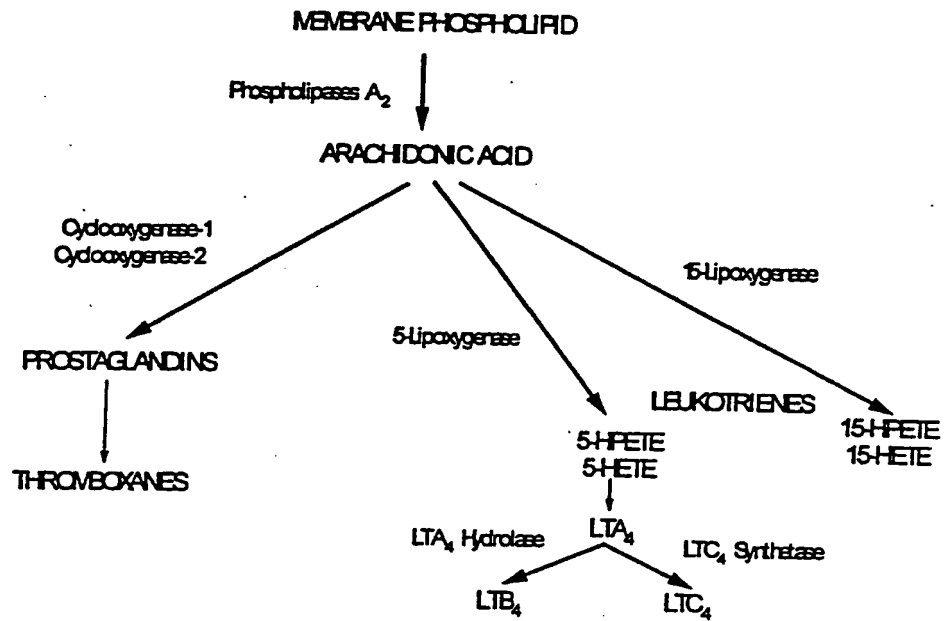


FIG. 3

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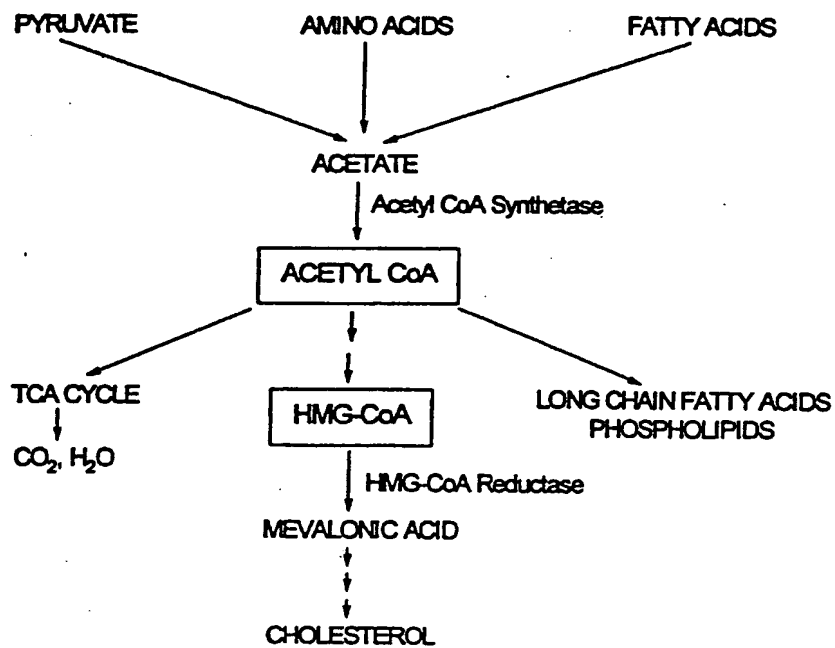


FIG. 4

SUBSTITUTE SHEET (RULE 26)



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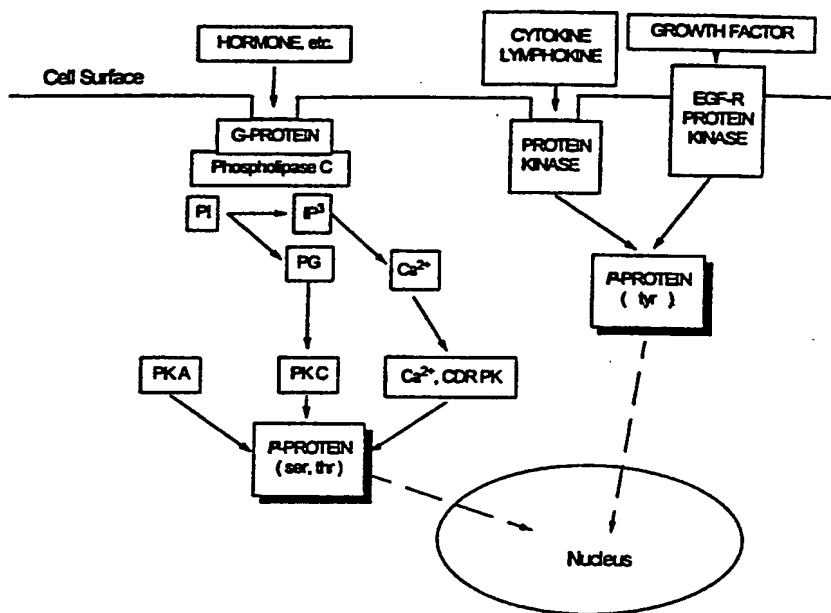


FIG. 5